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**(54) Title: SYNTHETIC PLANT GENES AND METHOD FOR PREPARATION****(57) Abstract**

A method for modifying structural gene sequences to enhance the expression of the protein product is disclosed. Also disclosed are novel structural genes which encode insecticidal proteins of *B.t.k.* HD-1, *B.t.k.* HD-73, *B.t. tenebrionis*, *B.t. entomocidus*, 2 protein of *B.t.k.* HD-1, and the coat protein of potato leaf roll virus.

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SYNTHETIC PLANT GENES AND METHOD FOR PREPARATION

BACKGROUND OF THE INVENTION

5

The present invention relates to genetic engineering and more particularly to plant transformation in which a plant is transformed to express a heterologous gene.

10 Although great progress has been made in recent years with respect to transgenic plants which express foreign proteins such as herbicide resistant enzymes and viral coat proteins, very little is known about the major factors affecting expression of foreign 15 genes in plants. Several potential factors could be responsible in varying degrees for the level of protein expression from a particular coding sequence. The level of a particular mRNA in the cell is certainly a critical factor.

20 The potential causes of low steady state levels of mRNA due to the nature of the coding sequence are many. First, full length RNA synthesis might not occur at a high frequency. This could, for example, be caused by the premature termination of RNA during transcription or due to unexpected mRNA processing 25 during transcription. Second, full length RNA could be produced but then processed (splicing, polyA addition) in the nucleus in a fashion that creates a nonfunctional mRNA. If the RNA is properly synthesized, terminated and polyadenylated, it then 30 can move to the cytoplasm for translation. In the

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cytoplasm, mRNAs have distinct half lives that are determined by their sequences and by the cell type in which they are expressed. Some RNAs are very short-lived and some are much more long-lived. In addition, there is an effect, whose magnitude is uncertain, of translational efficiency on mRNA half-life. In addition, every RNA molecule folds into a particular structure, or perhaps family of structures, which is determined by its sequence. The particular structure of any RNA might lead to greater or lesser stability in the cytoplasm. Structure per se is probably also a determinant of mRNA processing in the nucleus. Unfortunately, it is impossible to predict, and nearly impossible to determine, the structure of any RNA (except for tRNA) in vitro or in vivo. However, it is likely that dramatically changing the sequence of an RNA will have a large effect on its folded structure. It is likely that structure per se or particular structural features also have a role in determining RNA stability.

Some particular sequences and signals have been identified in RNAs that have the potential for having a specific effect on RNA stability. This section summarizes what is known about these sequences and signals. These identified sequences often are A+T rich, and thus are more likely to occur in an A+T rich coding sequence such as a B.t. gene. The sequence motif ATTAA (or AUUUA as it appears in RNA) has been implicated as a destabilizing sequence in mammalian cell mRNA (Shaw and Kamen, 1986). No analysis of the function of this sequence in plants has been done.

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Many short lived mRNAs have A+T rich 3' untranslated regions, and these regions often have the ATTAA sequence, sometimes present in multiple copies or as 5 multimers (e.g., ATTATATTAA...). Shaw and Kamen showed that the transfer of the 3' end of an unstable mRNA to a stable RNA (globin or VA1) decreased the stable RNA's half life dramatically. They further showed that a pentamer of ATTAA had a profound destabilizing 10 effect on a stable message, and that this signal could exert its effect whether it was located at the 3' end or within the coding sequence. However, the number of ATTAA sequences and/or the sequence context in which they occur also appear to be important in determining 15 whether they function as destabilizing sequences. Shaw and Kamen showed that a trimer of ATTAA had much less effect than a pentamer on mRNA stability and a dimer or a monomer had no effect on stability (Shaw and Kamen, 1987). Note that multimers of ATTAA such 20 as a pentamer automatically create an A+T rich region. This was shown to be a cytoplasmic effect, not nuclear. In other unstable mRNAs, the ATTAA sequence may be present in only a single copy, but it is often contained in an A+T rich region. From the animal cell data collected to date, it appears that ATTAA at least 25 in some contexts is important in stability, but it is not yet possible to predict which occurrences of ATTAA are destabilizing elements or whether any of these effects are likely to be seen in plants.

Some studies on mRNA degradation in animal cells 30 also indicate that RNA degradation may begin in some cases with nucleolytic attack in A+T rich regions. It

is not clear if these cleavages occur at ATTTA sequences. There are also examples of mRNAs that have differential stability depending on the cell type in which they are expressed or on the stage within the cell cycle at which they are expressed. For example, histone mRNAs are stable during DNA synthesis but unstable if DNA synthesis is disrupted. The 3' end of some histone mRNAs seems to be responsible for this effect (Pandey and Marzluff, 1987). It does not appear to be mediated by ATTTA, nor is it clear what controls the differential stability of this mRNA. Another example is the differential stability of IgG mRNA in B lymphocytes during B cell maturation (Genovese and Milcarek, 1988). A final example is the instability of a mutant beta-thalassemic globin mRNA. In bone marrow cells, where this gene is normally expressed, the mutant mRNA is unstable, while the wild-type mRNA is stable. When the mutant gene is expressed in HeLa or L cells in vitro, the mutant mRNA shows no instability (Lim et al., 1988). These examples all provide evidence that mRNA stability can be mediated by cell type or cell cycle specific factors. Furthermore this type of instability is not yet associated with specific sequences. Given these uncertainties, it is not possible to predict which RNAs are likely to be unstable in a given cell. In addition, even the ATTTA motif may act differentially depending on the nature of the cell in which the RNA is present. Shaw and Kamen (1987) have reported that activation of protein kinase C can block degradation mediated by ATTTA.

The addition of a polyadenylate string to the 3' end is common to most eucaryotic mRNAs, both plant and animal. The currently accepted view of polyA addition 5 is that the nascent transcript extends beyond the mature 3' terminus. Contained within this transcript are signals for polyadenylation and proper 3' end formation. This processing at the 3' end involves cleavage of the mRNA and addition of polyA to the 10 mature 3' end. By searching for consensus sequences near the polyA tract in both plant and animal mRNAs, it has been possible to identify consensus sequences that apparently are involved in polyA addition and 3' end cleavage. The same consensus sequences seem to be 15 important to both of these processes. These signals are typically a variation on the sequence AATAAA. In animal cells, some variants of this sequence that are functional have been identified; in plant cells there seems to be an extended range of functional sequences 20 (Wickens and Stephenson, 1984; Dean et al., 1986). Because all of these consensus sequences are variations on AATAAA, they all are A+T rich sequences. This sequence is typically found 15 to 20 bp before 25 the polyA tract in a mature mRNA. Experiments in animal cells indicate that this sequence is involved in both polyA addition and 3' maturation. Site directed mutations in this sequence can disrupt these functions (Conway and Wickens, 1988; Wickens et al., 1987). However, it has also been observed that 30 sequences up to 50 to 100 bp 3' to the putative polyA signal are also required; i.e., a gene that has a normal AATAAA but has been replaced or disrupted

downstream does not get properly polyadenylated (Gil and Proudfoot, 1984; Sadofsky and Alwine, 1984; McDevitt et al., 1984). That is, the polyA signal itself is not sufficient for complete and proper processing. It is not yet known what specific downstream sequences are required in addition to the polyA signal, or if there is a specific sequence that has this function. Therefore, sequence analysis can only identify potential polyA signals.

In naturally occurring mRNAs that are normally polyadenylated, it has been observed that disruption of this process, either by altering the polyA signal or other sequences in the mRNA, profound effects can be obtained in the level of functional mRNA. This has been observed in several naturally occurring mRNAs, with results that are gene specific so far. There are no general rules that can be derived yet from the study of mutants of these natural genes, and no rules that can be applied to heterologous genes. Below are four examples:

1. In a globin gene, absence of a proper polyA site leads to improper termination of transcription. It is likely, but not proven, that the improperly terminated RNA is nonfunctional and unstable (Proudfoot et al., 1987).
2. In a globin gene, absence of a functional polyA signal can lead to a 100-fold decrease in the level of mRNA accumulation (Proudfoot et al., 1987).
3. A globin gene polyA site was placed into the 3' ends of two different histone genes. The histone genes contain a secondary structure (stem-loop) near

their 3' ends. The amount of properly polyadenylated histone mRNA produced from these chimeras decreased as the distance between the stem-loop and the polyA site 5 increased. Also, the two histone genes produced greatly different levels of properly polyadenylated mRNA. This suggests an interaction between the polyA site and other sequences on the mRNA that can modulate mRNA accumulation (Pandy and Marzluff, 1987).

10 4. The soybean leghemoglobin gene has been cloned into HeLa cells, and it has been determined that this plant gene contains a "cryptic" polyadenylation signal that is active in animal cells, but is not utilized in plant cells. This leads to the production of a new 15 polyadenylated mRNA that is nonfunctional. This again shows that analysis of a gene in one cell type cannot predict its behavior in alternative cell types (Wiebauer et al., 1988).

20 From these examples, it is clear that in natural mRNAs proper polyadenylation is important in mRNA accumulation, and that disruption of this process can effect mRNA levels significantly. However, insufficient knowledge exists to predict the effect of changes in a normal gene. In a heterologous gene, 25 where we do not know if the putative polyA sites (consensus sequences) are functional, it is even harder to predict the consequences. However, it is possible that the putative sites identified are dysfunctional. That is, these sites may not act as proper polyA sites, but instead function as aberrant 30 sites that give rise to unstable mRNAs.

In animal cell systems, AATAAA is by far the most common signal identified in mRNAs upstream of the polyA, but at least four variants have also been found (Wickens and Stephenson, 1984). In plants, not nearly so much analysis has been done, but it is clear that multiple sequences similar to AATAAA can be used. The plant sites below called major or minor refer only to the study of Dean et al. (1986) which analyzed only three types of plant gene. The designation of polyadenylation sites as major or minor refers only to the frequency of their occurrence as functional sites in naturally occurring genes that have been analyzed. In the case of plants this is a very limited database. It is hard to predict with any certainty that a site designated major or minor is more or less likely to function partially or completely when found in a heterologous gene such as *B.t.*

20	PA	AATAAA	Major consensus site
	P1A	AATAAT	Major plant site
	P2A	AACCAA	Minor plant site
	P3A	ATATAA	"
	P4A	AATCAA	"
	P5A	ATACTA	"
25	P6A	ATAAAA	"
	P7A	ATGAAA	"
	P8A	AAGCAT	"
	P9A	ATTAAT	"
	P10A	ATACAT	"
30	P11A	AAAATA	"

	P12A	ATTA	AAA	Minor animal site
	P13A	AATT	AA	"
	P14A	AATA	CA	"
5	P15A	CATA	AA	"

Another type of RNA processing that occurs in the nucleus is intron splicing. Nearly all of the work on intron processing has been done in animal cells, but 10 some data is emerging from plants. Intron processing depends on proper 5' and 3' splice junction sequences. Consensus sequences for these junctions have been derived for both animal and plant mRNAs, but only a few nucleotides are known to be invariant. Therefore, 15 it is hard to predict with any certainty whether a putative splice junction is functional or partially functional based solely on sequence analysis. In particular, the only invariant nucleotides are GT at the 5' end of the intron and AG at the 3' end of the 20 intron. In plants, at every nearby position, either within the intron or in the exon flanking the intron, all four nucleotides can be found, although some positions show some nucleotide preference (Brown, 1986; Hanley and Schuler, 1988).

A plant intron has been moved from a patatin gene 25 into a GUS gene. To do this, site directed mutagenesis was performed to introduce new restriction sites, and this mutagenesis changed several nucleotides in the intron and exon sequences flanking the GT and AG. This intron still functioned properly, 30 indicating the importance of the GT and AG and the flexibility at other nucleotide positions. There are

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of course many occurrences of GT and AG in all genes that do not function as intron splice junctions, so there must be some other sequence or structural features that identify splice junctions. In plants, one such feature appears to be base composition per se. Wiebauer et al. (1988) and Goodall et al. (1988) have analyzed plant introns and exons and found that exons have ~50% A+T while introns have ~70% A+T. Goodall et al. (1988) also created an artificial plant intron that has consensus 5' and 3' splice junctions and a random A+T rich internal sequence. This intron was spliced correctly in plants. When the internal segment was replaced by a G+C rich sequence, splicing efficiency was drastically reduced. These two examples demonstrate that intron recognition in plants may depend on very general features -- splice junctions that have a great deal of sequence diversity and A+T richness of the intron itself. This, of course, makes it difficult to predict from sequence alone whether any particular sequence is likely to function as an active or partially active intron for RNA processing.

B.t. genes being A+T rich contain numerous stretches of various lengths that have 70% or greater A+T. The number of such stretches identified by sequence analysis depends on the length of sequence scanned.

As for polyadenylation described above, there are complications in predicting what sequences might be utilized as splice sites in any given gene. First, many naturally occurring genes have alternative

splicing pathways that create alternative combinations of exons in the final mRNA (Gallega and Nadal-Ginard, 1988; Helfman and Ricci, 1988; Tsurushita and Korn, 1989). That is, some splice junctions are apparently recognized under some circumstances or in certain cell types, but not in others. The rules governing this are not understood. In addition, there can be an interaction between processing paths such that utilization of a particular polyadenylation site can interfere with splicing at a nearby splice site and vice versa (Adami and Nevins, 1988; Brady and Wold, 1988; Marzluff and Pandey, 1988). Again no predictive rules are available. Also, sequence changes in a gene can drastically alter the utilization of particular splice junctions. For example, in a bovine growth hormone gene, small deletions in an exon a few hundred bases downstream of an intron cause the splicing efficiency of the intron to drop from greater than 95% to less than 2% (essentially nonfunctional). Other deletions however have essentially no effect (Hampson and Rottman, 1988). Finally, a variety of in vitro and in vivo experiments indicate that mutations that disrupt normal splicing lead to rapid degradation of the RNA in the nucleus. Splicing is a multistep process in the nucleus and mutations in normal splicing can lead to blockades in the process at a variety of steps. Any of these blockades can then lead to an abnormal and unstable RNA. Studies of mutants of normally processed (polyadenylation and splicing) genes are relevant to the study of heterologous genes such as B.t. B.t. genes might

contain functional signals that lead to the production of aberrant nonfunctional mRNAs, and these mRNAs are likely to be unstable. But the *B.t.* genes are perhaps even more likely to contain signals that are analogous to mutant signals in a natural gene. As shown above these mutant signals are very likely to cause defects in the processing pathways whose consequence is to produce unstable mRNAs.

It is not known with any certainty what signals RNA transcription termination in plant or animal cells. Some studies on animal genes that indicate that stretches of sequence rich in T cause termination by calf thymus RNA polymerase II in vitro. These studies have shown that the 3' ends of in vitro terminated transcripts often lie within runs of T such as T5, T6 or T7. Other identified sites have not been composed solely of T, but have had one or more other nucleotides as well. Termination has been found to occur within the sequences TATTTTTT, ATTCTC, TTCTT (Dedrick et al., 1987; Reines et al., 1987). In the case of these latter two, the context in which the sequence is found has been C+T rich as well. It is not known if this is essential. Other studies have implicated stretches of A as potential transcriptional terminators. An interesting example from SV40 illustrates the uncertainty in defining terminators based on sequence alone. One potential terminator in SV40 was identified as being A rich and having a region of dyad symmetry (potential stem-loop) 5' to the A rich stretch. However, a second terminator identified experimentally downstream in the same gene

5 was not A rich and included no potential secondary structure (Kessler et al., 1988). Of course, due to the A+T content of *B.t.* genes, they are rich in runs of A or T that could act as terminators. The importance of termination to stability of the mRNA is shown by the globin gene example described above. Absence of a normal polyA site leads to a failure in proper termination with a consequent decrease in mRNA.

10 There is also an effect on mRNA stability due to the translation of the mRNA. Premature translational termination in human triose phosphate isomerase leads to instability of the mRNA (Daar et al., 1988). Another example is the beta-thalassemic globin mRNA 15 described above that is specifically unstable in bone marrow cells (Lim et al., 1988). The defect in this mutant gene is a single base pair deletion at codon 44 that leads to translational termination (a nonsense codon) at codon 60. Compared to properly translated normal globin mRNA, this mutant RNA is very unstable. These results indicate that an improperly translated mRNA is unstable. Other work in yeast indicates that proper but poor translation can have an effect on mRNA levels. A heterologous gene was modified to convert 20 certain codons to more yeast preferred codons. An overall 10-fold increase in protein production was achieved, but there was also about a 3-fold increase in mRNA (Hoekema et al., 1987). This indicates that more efficient translation can lead to greater mRNA stability, and that the effect of codon usage can be 25 at the RNA level as well as the translational level. It is not clear from codon usage studies which codons 30

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lead to poor translation, or how this is coupled to mRNA stability.

5 Therefore, it is an object of the present invention to provide a method for preparing synthetic plant genes which express their respective proteins at relatively high levels when compared to wild-type genes. It is yet another object of the present invention to provide synthetic plant genes which 10 express the crystal protein toxin of *Bacillus thuringiensis* at relatively high levels.

BRIEF DESCRIPTION OF THE DRAWINGS

15 Figure 1 illustrates the steps employed in modifying a wild-type gene to increase expression efficiency in plants.

20 Figure 2 illustrates a comparison of the changes in the modified *B.t.k.* HD-1 sequence of Example 1 (lower line) versus the wild-type sequence of *B.t.k.* HD-1 which encodes the crystal protein toxin (upper line).

25 Figure 3 illustrates a comparison of the changes in the synthetic *B.t.k.* HD-1 sequence of Example 2 (lower line) versus the wild-type sequence of *B.t.k.* HD-1 which encodes the crystal protein toxin (upper line).

Figure 4 illustrates a comparison of the changes in the synthetic *B.t.k.* HD-73 sequence of Example 3 (lower line) versus the wild-type sequence of *B.t.k.* HD-73 (upper line).

30 Figure 5 represents a plasmid map of intermediate plant transformation vector cassette pMON893.

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Figure 6 represents a plasmid map of intermediate plant transformation vector cassette pMON900.

5 Figure 7 represents a map for the disarmed T-DNA of *A. tumefaciens* ACO.

Figure 8 illustrates a comparison of the changes in the synthetic truncated *B.t.k.* HD-73 gene (Amino acids 29-615 with an N-terminal Met-Ala) of Example 3 (lower line) versus the wild-type sequence of *B.t.k.* HD-73 10 (upper line).

Figure 9 illustrates a comparison of the changes in the synthetic/wild-type full length *B.t.k.* HD-73 sequence of Example 3 (lower line) versus the wild-type full-length sequence of *B.t.k.* HD-73 (upper 15 line).

Figure 10 illustrates a comparison of the changes in the synthetic/modified full length *B.t.k.* HD-73 sequence of Example 3 (lower line) versus the wild-type full-length sequence of *B.t.k.* HD-73 (upper 20 line).

Figure 11 illustrates a comparison of the changes in the fully synthetic full-length *B.t.k.* HD-73 sequence of Example 3 (lower line) versus the wild-type full-length sequence of *B.t.k.* HD-73 (upper 25 line).

Figure 12 illustrates a comparison of the changes in the synthetic *B.t.t.* sequence of Example 5 (lower line) versus the wild-type sequence of *B.t.t.* which encodes the crystal protein toxin (upper line).

Figure 13 illustrates a comparison of the changes 30 in the synthetic *B.t.* P2 sequence of Example 6 (lower

line) versus the wild-type sequence of *B.t.k.* HD-1 which encodes the P2 protein toxin (upper line).

5 Figure 14 illustrates a comparison of the changes in the synthetic *B.t.* *entomocidus* sequence of Example 7 (lower line) versus the wild-type sequence of *B.t.* *entomocidus* which encodes the Btent protein toxin (upper line).

10 Figure 15 illustrates a plasmid map for plant expression cassette vector pMON744.

Figure 16 illustrates a comparison of the changes in the synthetic potato leaf roll virus (PLRV) coat protein sequence of Example 9 (lower line) versus the wild-type coat protein sequence of PLRV (upper line).

15

#### STATEMENT OF THE INVENTION

20 The present invention provides a method for preparing synthetic plant genes which genes express their protein product at levels significantly higher than the wild-type genes which were commonly employed in plant transformation heretofore. In another aspect, the present invention also provides novel synthetic plant genes which encode non-plant proteins.

25 For brevity and clarity of description, the present invention will be primarily described with respect to the preparation of synthetic plant genes which encode the crystal protein toxin of *Bacillus thuringiensis* (*B.t.*). Suitable *B.t.* subspecies include, but are not limited to, *B.t. kurstaki* HD-1, *B.t. kurstaki* HD-73, 30 *B.t. sotto*, *B.t. berliner*, *B.t. thuringiensis*, *B.t. tolworthi*, *B.t. dendrolimus*, *B.t. alesti*, *B.t.*

*galleriae*, *B.t. aizawai*, *B.t. subtoxicus*, *B.t. entomocidus*, *B.t. tenebrionis* and *B.t. san diego*.

5 However, those skilled in the art will recognize and it should be understood that the present method may be used to prepare synthetic plant genes which encode non-plant proteins other than the crystal protein toxin of *B.t.* as well as plant proteins (see for instance, Example 9).

10 The expression of *B.t.* genes in plants is problematic. Although the expression of *B.t.* genes in plants at insecticidal levels has been reported, this accomplishment has not been straightforward. In particular, the expression of a full-length 15 lepidopteran specific *B.t.* gene (comprising DNA from a *B.t.k.* isolate) has been reported to be unsuccessful in yielding insecticidal levels of expression in some plant species (Vaeck et al., 1987 and Barton et al., 1987).

20 It has been reported that expression of the full-length gene from *B.t.k.* HD-1 was detectable in tomato plants but that truncated genes led to a higher frequency of insecticidal plants with an overall higher level of expression. Truncated genes of *B.t. berliner* also led to a higher frequency of 25 insecticidal plants in tobacco (Vaeck et al., 1987). On the other hand, insecticidal plants were provided from lettuce transformants using a full-length gene.

It has also been reported that the full length gene from *B.t.k.* HD-73 gave some insecticidal effect in 30 tobacco (Adang et al., 1987). However, the *B.t.* mRNA detected in these plants was only 1.7 kb compared to

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the expected 3.7 kb indicating improper expression of the gene. It was suggested that this truncated mRNA was too short to encode a functional truncated toxin, but there must have been a low level of longer mRNA in some plants or no insecticidal activity would have been observed. Others have reported in a publication that they observed a large amount of shorter than expected mRNA from a truncated *B.t.k.* gene, but some 5 mRNA of the expected size was also observed. In fact, it was suggested that expression of the full length gene is toxic to tobacco callus (Barton et al., 1987). The above illustrates that lepidopteran type *B.t.* genes are poorly expressed in plants compared to other 10 chimeric genes previously expressed from the same 15 promoter cassettes.

The expression of *B.t.t.* in tomato and potato is at levels similar to that of *B.t.k.* (i.e., poor). *B.t.t.* and *B.t.k.* genes share only limited sequence homology, 20 but they share many common features in terms of base composition and the presence of particular A+T rich elements.

All reports in the field have noted the lower than expected expression of *B.t.* genes in plants. In 25 general, insecticidal efficacy has been measured using insects very sensitive to *B.t.* toxin such as tobacco hornworm. Although it has been possible to obtain plants totally protected against tobacco hornworm, it is important to note that hornworm is up to 500 fold more sensitive to *B.t.* toxin than some agronomically 30 important insect pests such as beet armyworm. It is therefore of interest to obtain transgenic plants that

are protected against all important lepidopteran pests (or against Colorado potato beetle in the case of *B.t. tenebrionis*), and in addition to have a level of *B.t.* expression that provides an additional safety margin over and above the efficacious protection level. It is also important to devise plant genes which function reproducibly from species to species, so that insect resistant plants can be obtained in a predictable fashion.

In order to achieve these goals, it is important to understand the nature of the poorer than expected expression of *B.t.* genes in plants. The level of stable *B.t.* mRNA in plants is much lower than expected. That is, compared to other coding sequences driven by the same promoter, the level of *B.t.* mRNA measured by Northern analysis or nuclease protection experiments is much lower. For example, tomato plant 337 (Fischhoff et al., 1987) was selected as the best expressing plant with pMON9711 which contains the *B.t.k.* HD-1 *Kpn*I fragment driven by the CaMV 35S promoter and contains the NOS-NPTII-NOS selectable marker gene. In this plant the level of *B.t.* mRNA is between 100 to 1000 fold lower than the level of NPTII mRNA, even though the 35S promoter is approximately 50-fold stronger than the NOS promoter (Sanders et al., 1987).

The level of *B.t.* toxin protein detected in plants is consistent with the low level of *B.t.* mRNA. Moreover, the insecticidal efficacy of the transgenic plants correlates with the *B.t.* protein level indicating that the toxin protein produced in plants

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is biologically active. Therefore, the low level of *B.t.* toxin expression may be the result of the low levels of *B.t.* mRNA.

5       Messenger RNA levels are determined by the rate of synthesis and rate of degradation. It is the balance between these two that determines the steady state level of mRNA. The rate of synthesis has been maximized by the use of the CaMV 35S promoter, a  
10      strong constitutive plant expressible promoter. The use of other plant promoters such as nopaline synthase (NOS), mannopine synthase (MAS) and ribulose bisphosphatecarboxylase small subunit (RUBISCO) have not led to dramatic changes in the levels of *B.t.*  
15      toxin protein expression indicating that the effects determining *B.t.* toxin protein levels are promoter independent. These data imply that the coding sequences of DNA genes encoding *B.t.* toxin proteins are somehow responsible for the poor expression level, and that this effect is manifested by a low level of  
20      accumulated stable mRNA.

Lower than expected levels of mRNA have been observed with four different lepidopteran specific genes (two from *B.t.k.* HD-1; *B.t. berliner* and *B.t.k.* HD-73) as well as the gene from the coleopteran specific *B.t. tenebrionis*. It appears that for lepidopteran type *B.t.* genes these effects are manifest more strongly in the full length coding sequences than in the truncated coding sequences. These effects are seen across plant species although  
25      30      their magnitude seems greater in some plant species such as tobacco.

5        The nature of the coding sequences of *B.t.* genes distinguishes them from plant genes as well as many other heterologous genes expressed in plants. In particular, *B.t.* genes are very rich (~62%) in adenine (A) and thymine (T) while plant genes and most bacterial genes which have been expressed in plants are on the order of 45-55% A+T. The A+T content of the genomes (and thus the genes) of any organism are 10 features of that organism and reflect its evolutionary history. While within any one organism genes have similar A+T content, the A+T content can vary tremendously from organism to organism. For example, some *Bacillus* species have among the most A+T rich 15 genomes while some *Steptomyces* species are among the least A+T rich genomes (~30 to 35% A+T).

20        Due to the degeneracy of the genetic code and the limited number of codon choices for any amino acid, most of the "excess" A+T of the structural coding sequences of some *Bacillus* species are found in the third position of the codons. That is, genes of some *Bacillus* species have A or T as the third nucleotide in many codons. Thus A+T content in part can determine codon usage bias. In addition, it is clear 25 that genes evolve for maximum function in the organism in which they evolve. This means that particular nucleotide sequences found in a gene from one organism, where they may play no role except to code for a particular stretch of amino acids, have the potential to be recognized as gene control elements in 30 another organism (such as transcriptional promoters or terminators, polyA addition sites, intron splice

5 sites, or specific mRNA degradation signals). It is perhaps surprising that such misread signals are not a more common feature of heterologous gene expression,  
10 but this can be explained in part by the relatively homogeneous A+T content (~50%) of many organisms. This A+T content plus the nature of the genetic code put clear constraints on the likelihood of occurrence of any particular oligonucleotide sequence. Thus, a  
15 gene from *E. coli* with a 50% A+T content is much less likely to contain any particular A+T rich segment than a gene from *B. thuringiensis*.

15 As described above, the expression of *B.t.* toxin protein in plants has been problematic. Although the  
20 observations made in other systems described above offer the hope of a means to elevate the expression level of *B.t.* toxin proteins in plants, the success obtained by the present method is quite unexpected. Indeed, inasmuch as it has been recently reported that  
25 expression of the full-length *B.t.k.* toxin protein in tobacco makes callus tissue necrotic (Barton et al., 1987); one would reasonably expect that high level expression of *B.t.* toxin protein to be unattainable due to the reported toxicity effects.

25 In its most rigorous application, the method of the present invention involves the modification of an existing structural coding sequence ("structural gene") which codes for a particular protein by removal of ATTTA sequences and putative polyadenylation signals by site directed mutagenesis of the DNA  
30 comprising the structural gene. It is most preferred that substantially all the polyadenylation signals and

ATTTA sequences are removed although enhanced expression levels are observed with only partial removal of either of the above identified sequences.

5 Alternately if a synthetic gene is prepared which codes for the expression of the subject protein, codons are selected to avoid the ATTTA sequence and putative polyadenylation signals. For purposes of the present invention putative polyadenylation signals

10 include, but are not necessarily limited to, AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA. In replacing the ATTTA sequences and polyadenylation signals, codons are

15 preferably utilized which avoid the codons which are rarely found in plant genomes.

Another embodiment of the present invention, represented in the flow diagram of Figure 1, employs a method for the modification of an existing structural gene or alternately the *de novo* synthesis of a structural gene which method is somewhat less rigorous than the method first described above. Referring to Figure 1, the selected DNA sequence is scanned to identify regions with greater than four consecutive adenine (A) or thymine (T) nucleotides. The A+T

20 regions are scanned for potential plant polyadenylation signals. Although the absence of five or more consecutive A or T nucleotides eliminates most plant polyadenylation signals, if there are more than one of the minor polyadenylation signals identified

25 within ten nucleotides of each other, then the nucleotide sequence of this region is preferably

-24-

altered to remove these signals while maintaining the original encoded amino acid sequence.

5 The second step is to consider the 15 to 30 nucleotide regions surrounding the A+T rich region identified in step one. If the A+T content of the surrounding region is less than 80%, the region should be examined for polyadenylation signals. Alteration of the region based on polyadenylation signals is  
10 dependent upon (1) the number of polyadenylation signals present and (2) presence of a major plant polyadenylation signal.

15 The extended region is examined for the presence of plant polyadenylation signals. The polyadenylation signals are removed by site-directed mutagenesis of the DNA sequence. The extended region is also examined for multiple copies of the ATTAA sequence which are also removed by mutagenesis.

20 It is also preferred that regions comprising many consecutive A+T bases or G+C bases are disrupted since these regions are predicted to have a higher likelihood to form hairpin structure due to self-complementarity. Therefore, insertion of heterogeneous base pairs would reduce the likelihood of self-complementary secondary structure formation  
25 which are known to inhibit transcription and/or translation in some organisms. In most cases, the adverse effects may be minimized by using sequences which do not contain more than five consecutive A+T or G+C.

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SYNTHETIC OLIGONUCLEOTIDES FOR MUTAGENESIS

5        The oligonucleotides used in the mutagenesis are  
designed to maintain the proper amino acid sequence  
and reading frame and preferably to not introduce  
common restriction sites such as BglIII, HindIII, SacI,  
KpnI, EcoRI, NcoI, PstI and SalI into the modified  
gene. These restriction sites are found in multi-  
10      linker insertion sites of cloning vectors such as  
plasmids pUC118 and pMON7258. Of course, the  
introduction of new polyadenylation signals, ATTAA  
sequences or consecutive stretches of more than five  
A+T or G+C, should also be avoided. The preferred  
15      size for the oligonucleotides is around 40-50 bases,  
but fragments ranging from 18 to 100 bases have been  
utilized. In most cases, a minimum of 5 to 8 base  
pairs of homology to the template DNA on both ends of  
the synthesized fragment are maintained to insure  
proper hybridization of the primer to the template.  
20      The oligonucleotides should avoid sequences longer  
than five base pairs A+T or G+C. Codons used in the  
replacement of wild-type codons should preferably  
avoid the TA or CG doublet wherever possible. Codons  
are selected from a plant preferred codon table (such  
25      as Table I below) so as to avoid codons which are  
rarely found in plant genomes, and efforts should be  
made to select codons to preferably adjust the G+C  
content to about 50%.

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Table I

5

Preferred Codon Usage in Plants

10

<u>Amino Acid</u>	<u>Codon</u>	<u>Percent Usage in Plants</u>
ARG	CGA	7
	CGC	11
	CGG	5
	CGU	25
	AGA	29
	AGG	23

15

LEU	CUA	8
	CUC	20
	CUG	10
	CUU	28
	UUA	5
	UUG	30

20

SER	UCA	14
	UCC	26
	UCG	3
	UCU	21
	AGC	21
	AGU	15

THR	ACA	21
	ACC	41
	ACG	7
	ACU	31

25

PRO	CCA	45
	CCC	19
	CCG	9
	CCU	26

30

ALA	GCA	23
	GCC	32
	GCG	3
	GCU	41

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Table I - continued

Preferred Codon Usage in Plants

	<u>Amino Acid</u>	<u>Codon</u>	<u>Percent Usage in Plants</u>
5	GLY	GGA	32
		GGC	20
		GGG	11
		GGU	37
10	ILE	AUA	12
		AUC	45
		AUU	43
	VAL	GUA	9
15		GUC	20
		GUG	28
		GUU	43
	LYS	AAA	36
20	ASN	AAG	64
		AAC	72
	GLN	AAU	28
		CAA	64
25	HIS	CAG	36
		CAC	65
	GLU	CAU	35
		GAA	48
30	ASP	GAG	52
		GAC	48
	TYR	GAU	52
		UAC	68
	CYS	UAU	32
		UGC	78
		UGU	22

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Table I - continued

Preferred Codon Usage in Plants

5	<u>Amino Acid</u>	<u>Codon</u>	<u>Percent Usage in Plants</u>
10	PHE	UUC	56
		UUU	44
15	MET	AUG	100
		TRP	100

Regions with many consecutive A+T bases or G+C bases are predicted to have a higher likelihood to form hairpin structures due to self-complementarity. Disruption of these regions by the insertion of heterogeneous base pairs is preferred and should reduce the likelihood of the formation of self-complementary secondary structures such as hairpins which are known in some organisms to inhibit transcription (transcriptional terminators) and translation (attenuators). However, it is difficult to predict the biological effect of a potential hairpin forming region.

It is evident to those skilled in the art that while the above description is directed toward the modification of the DNA sequences of wild-type genes, the present method can be used to construct a completely synthetic gene for a given amino acid sequence. Regions with five or more consecutive A+T or G+C nucleotides should be avoided. Codons should be selected avoiding the TA and CG doublets in codons whenever possible. Codon usage can be normalized

against a plant preferred codon usage table (such as Table I) and the G+C content preferably adjusted to about 50%. The resulting sequence should be examined 5 to ensure that there are minimal putative plant polyadenylation signals and ATTAA sequences. Restriction sites found in commonly used cloning vectors are also preferably avoided. However, placement of several unique restriction sites 10 throughout the gene is useful for analysis of gene expression or construction of gene variants.

Plant Gene Construction

15 The expression of a plant gene which exists in double-stranded DNA form involves transcription of messenger RNA (mRNA) from one strand of the DNA by RNA polymerase enzyme, and the subsequent processing of the mRNA primary transcript inside the nucleus. This processing involves a 3' non-translated region which 20 adds polyadenylate nucleotides to the 3' end of the RNA. Transcription of DNA into mRNA is regulated by a region of DNA usually referred to as the "promoter." The promoter region contains a sequence of bases that signals RNA polymerase to associate with the DNA and 25 to initiate the transcription of mRNA using one of the DNA strands as a template to make a corresponding strand of RNA.

A number of promoters which are active in plant cells have been described in the literature. These 30 include the nopaline synthase (NOS) and octopine synthase (OCS) promoters (which are carried on tumor-

-30-

inducing plasmids of *Agrobacterium tumefaciens*), the Cauliflower Mosaic Virus (CaMV) 19S and 35S promoters, the light-inducible promoter from the small subunit of 5 ribulose bis-phosphate carboxylase (ssRUBISCO, a very abundant plant polypeptide) and the mannopine synthase (MAS) promoter (Velten et al. 1984 and Velten & Schell, 1985). All of these promoters have been used to create various types of DNA constructs which have 10 been expressed in plants (see e.g., PCT publication WO84/02913 (Rogers et al., Monsanto)).

Promoters which are known or are found to cause transcription of RNA in plant cells can be used in the present invention. Such promoters may be obtained 15 from plants or plant viruses and include, but are not limited to, the CaMV35S promoter and promoters isolated from plant genes such as ssRUBISCO genes. As described below, it is preferred that the particular promoter selected should be capable of causing 20 sufficient expression to result in the production of an effective amount of protein.

The promoters used in the DNA constructs (i.e. chimeric plant genes) of the present invention may be modified, if desired, to affect their control characteristics. For example, the CaMV35S promoter 25 may be ligated to the portion of the ssRUBISCO gene that represses the expression of ssRUBISCO in the absence of light, to create a promoter which is active in leaves but not in roots. The resulting chimeric promoter may be used as described herein. For 30 purposes of this description, the phrase "CaMV35S" promoter thus includes variations of CaMV35S promoter,

5 e.g., promoters derived by means of ligation with operator regions, random or controlled mutagenesis, etc. Furthermore, the promoters may be altered to contain multiple "enhancer sequences" to assist in elevating gene expression.

10 The RNA produced by a DNA construct of the present invention also contains a 5' non-translated leader sequence. This sequence can be derived from the promoter selected to express the gene, and can be 15 specifically modified so as to increase translation of the mRNA. The 5' non-translated regions can also be obtained from viral RNA's, from suitable eukaryotic genes, or from a synthetic gene sequence. The present invention is not limited to constructs, as presented 20 in the following examples. Rather, the non-translated leader sequence can be part of the 5' end of the non-translated region of the coding sequence for the virus coat protein, or part of the promoter sequence, or can be derived from an unrelated promoter or coding sequence. In any case, it is preferred that the sequence flanking the initiation site conform to the translational consensus sequence rules for enhanced 25 translation initiation reported by Kozak (1984).

25 The DNA construct of the present invention also contains a modified or fully-synthetic structural coding sequence which has been changed to enhance the performance of the gene in plants. In a particular embodiment of the present invention the enhancement 30 method has been applied to design modified and fully synthetic genes encoding the crystal toxin protein of *Bacillus thuringiensis*. The structural genes of the

present invention may optionally encode a fusion protein comprising an amino-terminal chloroplast transit peptide or secretory signal sequence (see for 5 instance, Examples 10 and 11).

The DNA construct also contains a 3' non-translated region. The 3' non-translated region contains a polyadenylation signal which functions in plants to cause the addition of polyadenylate nucleotides to the 10 3' end of the viral RNA. Examples of suitable 3' regions are (1) the 3' transcribed, non-translated regions containing the polyadenylation signal of *Agrobacterium* tumor-inducing (Ti) plasmid genes, such as the nopaline synthase (NOS) gene, and (2) plant 15 genes like the soybean storage protein (7S) genes and the small subunit of the RuBP carboxylase (E9) gene. An example of a preferred 3' region is that from the 7S gene, described in greater detail in the examples below.

20 Plant Transformation

A chimeric plant gene containing a structural coding sequence of the present invention can be inserted into the genome of a plant by any suitable 25 method. Suitable plants for use in the practice of the present invention include, but are not limited to, soybean, cotton, alfalfa, oilseed rape, flax, tomato, sugarbeet, sunflower, potato, tobacco, maize, rice and wheat. Suitable plant transformation vectors include 30 those derived from a Ti plasmid of *Agrobacterium tumefaciens*, as well as those disclosed, e.g., by

Herrera-Estrella (1983), Bevan (1983), Klee (1985) and EPO publication 120,516 (Schilperoort et al.). In addition to plant transformation vectors derived from the Ti or root-inducing (Ri) plasmids of *Agrobacterium*, alternative methods can be used to insert the DNA constructs of this invention into plant cells. Such methods may involve, for example, the use of liposomes, electroporation, chemicals that increase free DNA uptake, free DNA delivery via microprojectile bombardment, and transformation using viruses or pollen.

A particularly useful Ti plasmid cassette vector for transformation of dicotyledonous plants is shown in Figure 5. Referring to Figure 5, the expression cassette pMON893 consists of the enhanced CaMV35S promoter (EN 35S) and the 3' end including polyadenylation signals from a soybean gene encoding the alpha-prime subunit of beta-conglycinin. Between these two elements is a multilinker containing multiple restriction sites for the insertion of genes.

The enhanced CaMV35S promoter was constructed as follows. A fragment of the CaMV35S promoter extending between position -343 and +9 was previously constructed in pUC13 by Odell et al. (1985). This segment contains a region identified by Odell et al. (1985) as being necessary for maximal expression of the CaMV35S promoter. It was excised as a ClaI-HindIII fragment, made blunt ended with DNA polymerase I (Klenow fragment) and inserted into the HincII site of pUC18. This upstream region of the 35S promoter was excised from this plasmid as a HindIII-

5 EcoRV fragment (extending from -343 to -90) and inserted into the same plasmid between the HindIII and PstI sites. The enhanced CaMV35S promoter thus contains a duplication of sequences between -343 and -90 (Kay et al., 1987).

10 The 3' end of the 7S gene is derived from the 7S gene contained on the clone designated 17.1 (Schuler et al., 1982). This 3' end fragment, which includes the polyadenylation signals, extends from an Avall site located about 30 bp upstream of the termination codon for the beta-conglycinin gene in clone 17.1 to an EcoRI site located about 450 bp downstream of this termination codon.

15 The remainder of pMON893 contains a segment of pBR322 which provides an origin of replication in *E. coli* and a region for homologous recombination with the disarmed T-DNA in *Agrobacterium* strain ACO (described below); the oriV region from the broad host 20 range plasmid RK1; the streptomycin/spectinomycin resistance gene from Tn7; and a chimeric NPTII gene, containing the CaMV35S promoter and the nopaline synthase (NOS) 3' end, which provides kanamycin resistance in transformed plant cells.

25 Referring to Figure 6, transformation vector plasmid pMON900 is a derivative of pMON893. The enhanced CaMV35S promoter of pMON893 has been replaced with the 1.5kb mannopine synthase (MAS) promoter (Velten et al. 1984). The other segments are the same as plasmid pMON893. After incorporation of a DNA 30 construct into plasmid vector pMON893 or pMON900, the intermediate vector is introduced into *A. tumefaciens*

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strain ACO which contains a disarmed Ti plasmid. Cointegrate Ti plasmid vectors are selected and used to transform dicotyledonous plants.

5 Referring to Figure 7, *A. tumefaciens* ACO is a disarmed strain similar to pTiB6SE described by Fraley et al. (1985). For construction of ACO the starting *Agrobacterium* strain was the strain A208 which contains a nopaline-type Ti plasmid. The Ti plasmid  
10 was disarmed in a manner similar to that described by Fraley et al. (1985) so that essentially all of the native T-DNA was removed except for the left border and a few hundred base pairs of T-DNA inside the left border. The remainder of the T-DNA extending to a  
15 point just beyond the right border was replaced with a novel piece of DNA including (from left to right) a segment of pBR322, the oriV region from plasmid RK2, and the kanamycin resistance gene from Tn601. The pBR322 and oriV segments are similar to the segments  
20 in pMON893 and provide a region of homology for cointegrate formation.

The following examples are provided to better elucidate the practice of the present invention and should not be interpreted in any way to limit the scope of the present invention. Those skilled in the  
25 art will recognize that various modifications, truncations etc. can be made to the methods and genes described herein while not departing from the spirit and scope of the present invention.

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Example 1 -- Modified B.t.k. HD-1 Gene

5 Referring to Figure 2, the wild-type B.t.k. HD-1 gene is known to be expressed poorly in plants as a full length gene or as a truncated gene. The G+C content of the B.t.k. gene is low (37%) containing many A+T rich regions, potential polyadenylation sites (18 sites; see Table II for the list of sequences)  
10 and numerous ATTAA sequences.

Table II

15 List of Sequences of the Potential  
Polyadenylation Signals

	AATAAA*	AAGCAT
	AATAAT*	ATTAAT
	AACCAA	ATACAT
20	ATATAA	AAAATA
	AATCAA	ATTAAA**
	ATACTA	AATTAA**
	ATAAAAA	AATACA**
	ATGAAA	CATAAA**

25 \* indicates a potential major plant polyadenylation site.

\*\* indicates a potential minor animal polyadenylation site.

All others are potential minor plant polyadenylation sites.

30

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Table III lists the synthetic oligonucleotides designed and synthesized for the site-directed mutagenesis of the *B.t.k.* HD-1 gene.

5

Table III

Mutagenesis Primers for *B.t.k.* HD-1 Gene

10	<u>Primer</u>	<u>Length (bp)</u>	<u>Sequence</u>
	BTK185	18	TCCCCAGATA ATATCAAC
15	BTK240	48	GGCTTGATTCTAGCGAACT CTTCGATTCT CTGGTTGATG AGCTGTTC
20	BTK462	54	CAAAACTGAG AGGTGGAGGT TGGCAGCTTG AACGTACACG GAGAGGAGAGGAAC
25	BTK669	48	AGTTAGTGTA AGCTCTCTTC TGAACATGGTT GTACCTGATC CAATCTCT
30	BTK930	39	AGCCATGATC TGGTGACCGG ACCAGTAGTA TTCTCCTCT
	BTK1110	32	AGTTGTTGGT TGTTGATCCC GATGTTAAAA GG

Table III - continued

Mutagenesis Primers for B.t.k. HD-1 Gene

5

<u>Primer</u>	<u>Length (bp)</u>	<u>Sequence</u>
BTK1380A	37	GTGATGAAGG GATGATGTTG TTGAACTCAG CACTACG
BTK1380T	100	CAGAACGTTCC AGAGCCAAGA TTAGTAGACT TGGTGAGTGG GATTGGGTG ATTTGTGATG AAGGGATGAT GTTGTGAAC TCAGCACTAC GATGTATCCA
BTK1600	27	TGATGTGTGG AACTGAAGGT TTGTGGT

20        The *B.t.k.* HD-1 gene (BglII fragment from pMON9921 encoding amino acids 29-607 with a Met-Ala at the N-terminus) was cloned into pMON7258 (pUC118 derivative which contains a BglII site in the multilinker cloning region) at the BglII site resulting in pMON5342. The orientation of the *B.t.k.* gene was chosen so that the 25 opposite strand (negative strand) was synthesized in filamentous phage particles for the mutagenesis. The procedure of Kunkle (1985) was used for the mutagenesis using plasmid pMON5342 as starting material.

30        The regions for mutagenesis were selected in the following manner. All regions of the DNA sequence of

the *B.t.k.* gene were identified which contained five or more consecutive base pairs which were A or T. These were ranked in terms of length and highest percentage of A+T in the surrounding sequence over a 20-30 base pair region. The DNA was then analysed for regions which might contain polyadenylation sites (see Table II above) or ATTAA sequences. Oligonucleotides were designed which maximized the elimination of A+T consecutive regions which contained one or more polyadenylation sites or ATTAA sequences. Two potential plant polyadenylation sites were rated more critical (see Table II) based on published reports. Codons were selected which increased G+C content, did not generate restriction sites for enzymes useful for cloning and assembly of the modified gene (BamHI, BglIII, SacI, NcoI, EcoRV) and did not contain the doublets TA or GC which have been reported to be infrequently found in codons in plants. The oligonucleotides were at least 18 bp long ranging up to 100 base pairs and contained at least 5-8 base pairs of direct homology to native sequences at the ends of the fragments for efficient hybridization and priming in site-directed mutagenesis reactions. Figure 2 compares the wild-type *B.t.k.* HD-1 gene sequence with the sequence which resulted from the modifications by site-directed mutagenesis.

The end result of these changes was to increase the G+C content of *B.t.k.* gene from 37% to 41% while also decreasing the potential plant polyadenylation sites from 18 to 7 and decreasing the ATTAA regions from 13 to 7. Specifically, the mutagenesis changes from

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amino (5') terminus to the carboxy (3') terminus are as follows:

5 BTK185 is an 18-mer used to eliminate a plant polyadenylation site in the midst of a nine base pair region of A+T.

10 BTK240 is a 48-mer. Seven base pairs were changed by this oligonucleotide to eliminate three potential polyadenylation sites (2 AACCAA, 1 AATTAA). Another 15 region close to the region altered by BTK240, starting at bp 312, had a high A+T content (13 of 15 base pairs) and an ATTTA region. However, it did not contain a potential polyadenylation site and its longest string of uninterrupted A+T was seven base pairs.

15 BTK462 is a 54-mer introducing 13 base pair changes. The first six changes were to reduce the A+T richness of the gene by replacing wild-type codons with codons containing G and C while avoiding the CG doublet. The next seven changes made by BTK462 were 20 used to eliminate an A+T rich region (13 of 14 base pairs were A or T) containing two ATTTA regions.

25 BTK669 is a 48-mer making nine individual base pair changes eliminating three possible polyadenylation sites (ATATAA, AATCAA, and AATTAA) and a single ATTTA site.

30 BTK930 is a 39-mer designed to increase the G+C content and to eliminate a potential polyadenylation site (AATAAT - a major site). This region did contain a nine base pair region of consecutive A+T sequence. One of the base pair changes was a G to A because a G at this position would have created a G+C rich region

5 (CCGG(G)C). Since sequencing reactions indicate that there can be difficulties generating sequence through G+C consecutive bases, it was thought to be prudent to avoid generating potentially problematic regions even if they were problematic only in vitro.

10 BTK1110 is a 32-mer designed to introduce five changes in the wild-type gene. One potential site (AATAAT - a major site) was eliminated in the midst of an A+T rich region (19 of 22 base pairs).

15 BTK1380A and BTK1380T are responsible for 14 individual base pair changes. The first region (1380A) has 17 consecutive A+T base pairs. In this region is an ATTAA and a potential polyadenylation site (AATAAT). The 100-mer (1380T) contains all the changes dictated by 1380A. The large size of this primer was in part an experiment to determine if it was feasible to utilize large oligonucleotides for mutagenesis (over 60 bases in length). A second consideration was that the 100-mer was used to 20 mutagenize a template which had previously been mutagenized by 1380A. The original primer ordered to mutagenize the region downstream and adjacent to 1380A did not anneal efficiently to the desired site as indicated by an inability to obtain clean sequence 25 utilizing the primer. The large region of homology of 1380T did assure proper annealing. The extended size of 1380T was more of a convenience rather than a necessity. The second region adjacent to 1380A covered by 1380T has a high A+T content (22 of 29 30 bases are A or T).

5 BTK1600 is a 27-mer responsible for five individual base pair changes. An ATTTA region and a plant polyadenylation site were identified and the appropriate changes engineered.

10 A total of 62 bases were changed by site-directed mutagenesis. The G+C content increased by 55 base pairs, the potential polyadenylation sites were reduced from 18 to seven and the ATTTA sequences 15 decreased from 13 to seven. The changes in the DNA sequence resulted in changes in 55 of the 579 codons in the truncated *B.t.k.* gene in pMON5342 (approximately 9.5%).

15 Referring to Table IV modified *B.t.k.* HD-1 genes were constructed that contained all of the above modifications (pMON5370) or various subsets of individual modifications. These genes were inserted into pMON893 for plant transformation and tobacco plants containing these genes were analyzed. The 20 analysis of tobacco plants with the individual modifications was undertaken for several reasons. Expression of the wild type truncated gene in tobacco is very poor, resulting in infrequent identification of plants toxic to THW. Toxicity is defined by leaf feeding assays as at least 60% mortality of tobacco 25 hornworm neonate larvae with a damage rating of 1 or less (scale is 0 to 4; 0 is equivalent to total protection, 4 total damage). The modified HD-1 gene (pMON5370) shows a large increase in expression (estimated to be approximately 100-fold; see Table 30 VIII) in tobacco. Therefore, increases in expression of the wild-type gene due to individual modifications

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would be apparently a large increase in the frequency  
of toxic tobacco plants and the presence of detectable  
B.t.k. protein. Results are shown in the following  
5 table:

Table IV

10 Relative effects of Regional Modifications  
within the B.t.k. Gene

	<u>Construct</u>	<u>Position Modified</u>	<u># of Plants</u>	<u># of Toxic Plants</u>
15	PMON5370	185,240,669,930, 1110,1380a+b,1600	38	22
	PMON10707	185,240,462,669	48	19
20	PMON10706	930,1110,1380a+b,1600	43	1
	PMON10539	185	55	2
	PMON10537	240	57	17
25	PMON10540	185,240	88	23
	PMON10705	462	47	1

The effects of each individual oligonucleotides' changes on expression did reveal some overall trends.  
30 Six different constructs were generated which were

designed to identify the key regions. The nine different oligonucleotides were divided in half by their position on the gene. Changes in the N-terminal half were incorporated into pMON10707 (185,240, 462,669). C-terminal half changes were incorporated into pMON10706 (930,1110,1380a+b,1600). The results of analysis of plants with these two constructs indicate that pMON10707 produces a substantial number of toxic plants (19 of 48). Protein from these plants is detectable by ELISA analysis. pMON10706 plants were rarely identified as insecticidal (1 of 43) and the levels of *B.t.k.* were barely detectable by immunological analysis. Investigation of the N-terminal changes in greater detail was done with 4 pMON constructs; 10539 (185 alone), 10537 (240 alone), 10540 (185 and 240) and 10705 (462 alone). The results indicate that the presence of the changes in 240 were required to generate a substantial number of toxic plants (pMON10540; 23 of 88, pMON10537; 17 of 57). The absence of the 240 changes resulted in a low frequency of toxic plants with low *B.t.k.* protein levels, identical to results with the wild type gene. These results indicate that the changes in 240 are responsible for a substantial increase in *B.t.k.* expression levels over an analogous wild-type construct in tobacco. Changes in additional regions (185,462,669) in conjunction with 240 may result in increases in *B.t.k.* expression (>2 fold). However, changes at the 240 region of the N-terminal portion of the gene do result in dramatic increases in expression.

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Despite the importance of the alteration of the 240 region in expression of modified genes, increased expression can be achieved by alteration of other regions. Hybrid genes, part wild-type, part synthetic, were generated to determine the effects of synthetic gene segments on the levels of *B.t.k.* expression. A hybrid gene was generated with a synthetic N-terminal third (base pair 1 to 590 of Figure 2: to the *Xba*I site) with the C-terminal wild type *B.t.k.* HD-1 (pMON5378) Plants transformed with this vector were as toxic as plants transformed with the modified HD-1 gene (pMON5370). This is consistent with the alteration of the 240 region. However, pMON10538, a hybrid with a wild-type N-terminal third (wild type gene for the first 600 base pairs, to the second *Xba*I site) and a synthetic C-terminal last two-thirds (base pair 590 to 1845 of Figure 3 was used to transform tobacco and resulted in a dramatic increase in expression. The levels of expression do not appear to be as high as those seen with the synthetic gene, but are comparable to the modified gene levels. These results indicate that modification of the 240 segment is not essential to increased expression since pMON10538 has an intact 240 region. A fully synthetic gene is, in most cases, superior for expression levels of *B.t.k.* (See Example 2.)

Example 2 -- Fully Synthetic *B.t.k.* HD-1 Gene

A synthetic *B.t.k.* HD-1 gene was designed using the preferred plant codons listed in Table V below.

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Table V lists the codons and frequency of use in plant genes of dicotyledonous plants compared to the frequency of their use in the wild type *B.t.k.* HD-1 gene (amino acids 1-615) and the synthetic gene of this example. The total number of each amino acid in this segment of the gene is listed in the parenthesis under the amino acid designated.

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Table V

Codon in Usage Synthetic B.t.k. HD-1 Gene

5

			Percent Usage in		
			Plants	Wt. B.t.k.	/Syn
10	5 ARG (43)	CGA	7	11	2
		CGC	11	5	5
		CGG	5	2	0
		CGU	25	14	27
		AGA	29	55	41
		AGG	23	14	25
15	15 LEU (49)	CUA	8	16	4
		CUC	20	0	20
		CUG	10	2	6
		CUU	28	22	24
		UUA	5	50	0
		UUG	30	10	45
20	20 SER (64)	UCA	14	27	5
		UCC	26	9	28
		UCG	3	8	0
		UCU	21	19	31
		AGC	21	6	32
		AGU	15	31	5

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Table V - continued

5                   Codon in Usage Synthetic B.t.k. HD-1 Gene

	<u>Amino Acid</u>	<u>Codon</u>	<u>Percent Usage in Plants/Wt B.t.k./Syn</u>		
10	THR (42)	ACA	21	31	14
		ACC	41	19	53
		ACG	7	14	0
		ACU	31	36	33
15	PRO (34)	CCA	45	35	53
		CCC	19	6	12
		CCG	9	21	3
		CCU	26	38	32
20	ALA (31)	GCA	23	38	26
		GCC	32	9	29
		GCG	3	3	0
		GCU	41	50	45
25	GLY (46)	GGA	32	52	45
		GGC	20	17	15
		GGG	11	15	6
		GGU	37	15	34
30	ILE (46)	AUA	12	39	2
		AUC	45	11	67
		AUU	43	50	30

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Table V - continued

5      Codon in Usage Synthetic B.t.k. HD-1 Gene

	<u>Amino Acid</u>	<u>Codon</u>	Percent Usage in Plants/Wt B.t.k./Syn		
10	VAL (38)	GUA	9	45	3
		GUC	20	5	16
		GUG	28	11	37
		GUU	43	39	45
15	LYS (3)	AAA	36	100	33
		AAG	64	0	67
20	ASN (44)	AAC	72	27	80
		AAU	28	73	20
25	GLN (31)	CAA	64	77	61
		CAG	36	23	39
30	HIS (10)	CAC	65	0	80
		CAU	35	100	20
25	GLU (30)	GAA	48	87	50
		GAG	52	13	50
30	ASP (23)	GAC	48	17	65
		GAU	52	83	35

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5

Table V - continued

Codon in Usage Synthetic B.t.k. HD-1 Gene

	<u>Amino Acid</u>	<u>Codon</u>	Percent Usage in Plants/Wt B.t.k./Syn		
10	TYR	UAC	68	20	72
	(25)	UAU	32	80	28
15	CYS	UGC	78	50	100
	(2)	UGU	22	50	0
	PHE	UUC	56	17	83
	(36)	UUU	44	83	17
20	MET	AUG	100	100	100
	(9)				
	TRP	UGG	100	100	100
	(9)				

25

The resulting synthetic gene lacks ATTAA sequences, contains only one potential polyadenylation site and has a G+C content of 48.5%. Figure 3 is a comparison of the wild-type HD-1 sequence to the synthetic gene sequence for amino acids 1-615. There is approximately 77% DNA homology between the synthetic

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gene and the wild-type gene and 356 of the 615 codons have been changed (approximately 60%).

5

Example 3 -- Synthetic B.t.k. HD-73 Gene

The crystal protein toxin from B.t.k. HD-73 exhibits a higher unit activity against some important agricultural pests. The toxin protein of HD-1 and HD-10 73 exhibit substantial homology (~90%) in the N-terminal 450 amino acids, but differ substantially in the amino acid region 451-615. Fusion proteins comprising amino acids 1-450 of HD-1 and 451-615 of HD-15 73 exhibit the insecticidal properties of the wild-type HD-73. The strategy employed was to use the 5'- two thirds of the synthetic HD-1 gene (first 1350 bases, up to the SacI site) and to dramatically modify the final 590 bases (through amino acid 645) of the HD-20 73 in a manner consistent with the algorithm used to design the synthetic HD-1 gene. Table VI below lists the oligonucleotides used to modify the HD-73 gene in the order used in the gene from 5' to 3' end. Nine oligonucleotides were used in a 590 base pair region, each nucleotide ranging in size from 33 to 60 bases. The only regions left unchanged were areas where there 25 were no long consecutive strings of A or T bases (longer than six). All polyadenylation sites and ATTTA sites were eliminated.

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Table VI

<u>Mutagenesis Primers for B.t.k. HD-73</u>			
	<u>Primer</u>	<u>Length (bp)</u>	<u>Sequence</u>
5	73K1363	51	AATACTATCG GATGCGATGA TGTGTTGAA CTCAGCACTA CGGTGTATCC A
10	73K1437	33	TCCTGAAATG ACAGAACCGT TGAAGAGAAA GTT
15	73K1471	48	ATTTCCACTG CTGTTGAGTC TAACGAGGTC TCCACCAGTG AATCCTGG
20	73K1561	60	GTGAATAGGG GTCACAGAAG CATACTCAC ACGAACTCTA TATCTGGTAG ATGTTGGATGG
25	73K1642	33	TGTAGCTGGA ACTGTATTGG AGAAGATGGA TGA
30	73K1675	48	TTCAAAGTAA CCGAAATCGC TGGATTGGAG ATTATCCAAG GAGGTAGC
	73K1741	39	ACTAAAGTTT CTAACACCCA CGATGTTACC GAGTGAAGA

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Table VI - continued

Mutagenesis Primers for B.t.k. HD-73

5

<u>Primer</u>	<u>Length (bp)</u>	<u>Sequence</u>
73K1797	36	AACTGGAATG AACTCGAATC TGTCGATAAT CACTCC
73KTERM	54	GGACACTAGA TCTTAGTGAT AATCGGTACAC ATTTGTCTTG AGTCCAAGCT GGTT

10        15        The resulting gene has two potential polyadenylation sites (compared to 18 in the WT) and no ATTTA sequence (12 in the WT). The G+C content has increased from 37% to 48%. A total of 59 individual base pair changes were made using the primers in Table VI. Overall, there is 90% DNA homology between the region of the HD-73 gene modified by site directed mutagenesis and the wild-type sequence of the analogous region of HD-73. The synthetic HD-73 is a hybrid of the first 1360 bases from the synthetic HD-1 and the next 590 bases or so modified HD-73 sequence.

15        20        25        Figure 4 is a comparison of the above-described synthetic B.t.k. HD-73 and the wild-type B.t.k. HD-73 encoding amino acids 1-645. In the modified region of the HD-73 gene 44 of the 170 codons (25%) were changed as a result of the site-directed mutagenesis changes

20        25        30        resulting from the oligonucleotides found in Table VI. Overall, approximately 50% of the codons in the

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synthetic *B.t.k.* HD-73 differ from the analogous segment of the wild-type and HD-73 gene.

5 A one base pair deletion in the synthetic HD-73 gene was detected in the course of sequencing the 3' end at base pair 1890. This results in a frame-shift mutation at amino acid 625 with a premature stop codon at amino acid 640 (pMON5379). Table VII below compares the codon usage of the wild-type gene of *B.t.k.* HD-73 10 versus the synthetic gene of this example for amino acids 451-645 and codon usage of naturally occurring genes of dicotyledonous plants. The total number of each amino acid encoded in this segment of the gene is found in the parentheses under the amino acid 15 designation.

Table VII  
Codon Usage in Synthetic *B.t.k.* HD-73 Gene

	<u>Amino Acid</u>	<u>Codon</u>	<u>Percent Usage in Plants/Wt HD-73/Syn</u>		
20	ARG	CGA	7	10	0
	(10)	CGC	11	0	8
		CGG	5	10	0
		CGU	25	20	23
		AGA	29	60	62
		AGG	23	0	8

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Table VII - continued  
Codon Usage in Synthetic B.t.k. HD-73 Gene

		<u>Amino Acid</u>	<u>Codon</u>	<u>Percent Usage in</u>		
				<u>Plants/Wt HD-73/Syn</u>		
5	LEU	CUA	8	25	8	
		CUC	20	17	58	
		CUG	10	17	8	
	(12)	CUU	28	8	0	
		UUA	5	33	8	
		UUG	30	0	17	
10	SER	UCA	14	24	18	
		UCC	26	10	27	
		UCG	3	10	0	
		UCU	21	24	18	
		AGC	21	0	14	
		AGU	15	33	23	
15	THR	ACA	21	47	38	
		ACC	41	13	31	
		ACG	7	13	0	
		ACU	31	27	31	
20	PRO	CCA	45	71	71	
		CCC	19	0	0	
		CCG	9	14	0	
		CCU	26	14	29	
25						
30						

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Table VII - continued  
Codon Usage in Synthetic B.t.k. HD-73 Gene

			Percent Usage in Plants/Wt HD-73/Syn		
	<u>Amino Acid</u>	<u>Codon</u>	23	29	31
5	ALA	GCA	23	29	31
	(14)	GCC	32	7	8
		GCG	3	21	15
10		GCU	41	43	46
	GLY	GGA	32	33	43
	(15)	GGC	20	0	0
		GGG	11	27	14
15		GGU	37	40	43
	ILE	AUA	12	33	7
	(15)	AUC	45	7	40
		AUU	43	60	53
20	VAL	GUA	9	40	7
	(15)	GUC	20	0	7
		GUG	28	20	36
		GUU	43	40	50
25	LYS	AAA	36	67	100
	(3)	AAG	64	33	0
30	ASN	AAC	72	20	53
	(20)	AAU	28	80	47

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Table VII - continued  
Codon Usage in Synthetic B.t.k. HD-73 Gene

			Percent Usage in Plants/Wt HD-73/Syn		
	<u>Amino Acid</u>	<u>Codon</u>			
5	GLN	CAA	64	60	67
	(5)	CAG	36	40	33
10	HIS	CAC	65	67	100
	(3)	CAU	35	33	0
15	GLU	GAA	48	86	57
	(7)	GAG	52	14	43
20	ASP	GAC	48	40	50
	(5)	GAU	52	60	50
25	TYR	UAC	68	0	20
	(5)	UAU	32	100	80
30	CYS	UGC	78	0	0
	(0)	UGU	22	0	0
35	PHE	UUC	56	8	67
	(13)	UUU	44	92	33
40	MET	AUG	100	100	100
	(2)				
45	TRP	UGG	100	100	100
	(2)				

5        Another truncated synthetic HD-73 gene was constructed. The sequence of this synthetic HD-73 gene is identical to that of the above synthetic HD-73 gene in the region in which they overlap (amino acids 29-615), and it also encodes Met-Ala at the N-terminus. Figure 8 shows a comparison of this truncated synthetic HD-73 gene with the N-terminal Met-Ala versus the wild-type HD-73 gene.

10      While the previous examples have been directed at the preparation of synthetic and modified genes encoding truncated *B.t.k.* proteins, synthetic or modified genes can also be prepared which encode full length toxin proteins.

15      One full length *B.t.k.* gene consists of the synthetic HD-73 sequence of Figure 4 from nucleotide 1-1845 plus wild-type HD-73 sequence encoding amino acids 616 to the C-terminus of the native protein. Figure 9 shows a comparison of this synthetic/wild-type full length HD-73 gene versus the wild-type full length HD-73 gene.

20      Another full length *B.t.k.* gene consists of the synthetic HD-73 sequence of Figure 4 from nucleotide 1-1845 plus a modified HD-73 sequence ending amino acids 616 to the C-terminus of the native protein. The C-terminal portion has been modified by site-directed mutagenesis to remove putative polyadenylation signals and ATTTA sequences according to the algorithm of Figure 1. Figure 10 shows a comparison of this synthetic/modified full length HD-73 gene versus the wild-type full length HD-73 gene.

Another full length *B.t.k.* gene consists of a fully synthetic HD-73 sequence which incorporates the synthetic HD-73 sequence of Figure 4 from nucleotide 1-  
5 1845 plus a synthetic sequence encoding amino acids 616 to the C-terminus of the native protein. The C-terminal synthetic portion has been designed to eliminate putative polyadenylation signals and ATTAA sequences and to include plant preferred codons.  
10 Figure 11 shows a comparison of this fully synthetic full length HD-73 gene versus the wild-type full length HD-73 gene.

Alternatively, another full length *B.t.k.* gene consists of a fully synthetic sequence comprising base pairs 1-1830 of *B.t.k.* HD-1 (Figure 3) and base pairs 15 1834-3534 of *B.t.k.* HD-73 (Figure 11).

Example 4 -- Expression of Modified and Synthetic  
*B.t.k.* HD-1 and Synthetic HD-73

20 A number of plant transformation vectors for the expression of *B.t.k.* genes were constructed by incorporating the structural coding sequences of the previously described genes into plant transformation cassette vector pMON893. The respective intermediate  
25 transformation vector is inserted into a suitable disarmed *Agrobacterium* vector such as *A. tumefaciens* ACO, supra. Tissue explants are cocultured with the disarmed *Agrobacterium* vector and plants regenerated under selection for kanamycin resistance using known  
30 protocols: tobacco (Horsch et al., 1985); tomato

(McCormick et al., 1986) and cotton (Trolinder et al., 1987).

5 a) Tobacco.

The level of *B.t.k.* HD-1 protein in transgenic tobacco plants containing pMON9921 (wild type truncated), pMON5370 (modified HD-1, Example 1, Figure 2) and pMON5377 (synthetic HD-1, Example 2, Figure 3) 10 were analyzed by Western analysis. Leaf tissue was frozen in liquid nitrogen, ground to a fine powder and then ground in a 1:2 (wt:volume) of SDS-PAGE sample buffer. Samples were frozen on dry ice, then 15 incubated for 10 minutes in a boiling water bath and microfuged for 10 minutes. The protein concentration of the supernatant was determined by the method of Bradford (Anal. Biochem. 72:248-254). Fifty ug of protein was run per lane on 9% SDS-PAGE gels, the protein transferred to nitrocellulose and the *B.t.k.* 20 HD-1 protein visualized using antibodies produced against *B.t.k.* HD-1 protein as the primary antibody and alkaline phosphatase conjugated second antibody as described by the manufacturer (Promega, Madison, WI). Purified HD-1 tryptic fragment was used as the 25 control. Whereas the *B.t.k.* protein from tobacco plants containing pMON9921 was below the level of detection, the *B.t.k.* protein from plants containing the modified (pMON5370) and synthetic (pMON5377) genes was easily detected. The *B.t.k.* protein from plants containing pMON9921 remained undetectable, even with 30 10 fold longer incubation times. The relative levels of *B.t.k.* HD-1 protein in these plants is estimated in

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Table VIII. Because the protein from plants containing pMON9921 was not observed, the level of protein in these plants was estimated from the relative mRNA levels (see below). Plants containing the modified gene (pMON5370) expressed approximately 100 fold more *B.t.k.* protein than plants containing the wild-type gene (pMON9921). Plants containing the fully synthetic *B.t.k.* HD-1 gene (pMON5377) expressed approximately five fold more protein than plants containing the modified gene. The modified gene contributes the majority of the increase in *B.t.k.* expression observed. The plants used to generate the above data are the best representatives from each construct based either on a tobacco hornworm bioassay or on data derived from previous Western analysis.

Table VIII

Expression of *B.t.k.* HD-1 Protein  
in Transgenic Tobacco

<u>Gene Description</u>	<u>Vector</u>	<u><i>B.t.k.</i> Protein* Concentration</u>	<u>Fold Increase in <i>B.t.k.</i> Expression</u>
Wild type	pMON9921	10	1
Modified	pMON5370	1000	100
Synthetic	pMON5377	5000	500

\* *B.t.k.* protein concentrations are expressed in  
30 ng/mg of total soluble protein. The level of *B.t.k.*

protein for plants containing the wild type gene are estimated from mRNA levels.

5 Plants containing these genes were tested for bioactivity to determine whether the increased quantities of protein observed by Western analysis result in a corresponding increase in bioactivity. Leaves from the same plants used for the Western data  
10 in Table 1 were tested for bioactivity against two insects. A detached leaf bioassay was first done using tobacco hornworm, an extremely sensitive lepidopteran insect. Leaves from all three transgenic tobacco plants were totally protected and 100%  
15 mortality of tobacco hornworm observed (see Table IX below). A much less sensitive insect, beet armyworm, was then used in another detached leaf bioassay. Beet armyworm is approximately 500 fold less sensitive to *B.t.k.* HD-1 protein than tobacco hornworm. The  
20 difference in sensitivity of these two insects was determined using purified HD-1 protein in a diet incorporation assay (see below). Plants containing the wild-type gene (pMON9921) showed only minimal protection against beet armyworm, whereas plants containing the modified gene showed almost complete  
25 protection and plants containing the fully synthetic gene were totally protected against beet armyworm damage. The results of these bioassays confirm the levels of *B.t.k.* HD-1 expression observed in the Western analysis and demonstrates that the increased  
30 levels of *B.t.k.* HD-1 protein correlates with increased insecticidal activity.

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Table IX

5

Protection of Tobacco Plants from  
Tobacco Hornworm and Beet Armyworm

10

Gene		Tobacco Hornworm	Beet Armyworm
<u>Description</u>	<u>Vector</u>	<u>Damage*</u>	<u>Damage*</u>
None	None	NL	NL
Wild type	pMON9921	0	3
Modified	pMON5370	0	1
Synthetic	pMON5377	0	0

15

\* Extent of insect damage was rated: 0, no damage; 1, slight; 2, moderate; 3, severe; or NL, no leaf left.

20

The bioactivity of the *B.t.k.* HD-1 protein produced by these transgenic plants was further investigated to more accurately quantitate the relative activities. Leaf tissue from tobacco plants containing the wild-type, modified and synthetic genes were ground in 100 mM sodium carbonate buffer, pH 10 at a 1:2 (wt:vol) ratio. Particulate material was removed by centrifugation. The supernatant was incorporated into a synthetic diet similar to that described by Marrone et al. (1985). The diet medium was prepared the day of the test with the plant extract solutions incorporated in place of the 20% water component. One ml of the diet was aliquoted into 96 well plates.

After the diet dried, one neonate tobacco budworm larva was added to each well. Sixteen insects were tested with each plant sample. The plants were 5 incubated at 27°C. After seven days, the larvae from each treatment were combined and weighed on an analytical balance. The average weight per insect was calculated and compared to a standard curve relating B.t.k. protein concentrations to average larval 10 weight. Insect weight was inversely proportional (in a logarithmic manner) to the relative increase in B.t.k. protein concentration. The amount of B.t.k. HD-1 protein, based on the extent of larval growth inhibition was determined for two different plants 15 containing each of the three genes. The specific activity (ng of B.t.k. HD-1 per mg of plant protein) was determined for each plant. Plants containing the modified HD-1 gene (pMON5370) averaged approximately 1400 ng (1200 and 1600 ng) of B.t.k. HD-1 per mg of 20 plant extract protein. This value compares closely with the 1000 ng of B.t.k. HD-1 protein per mg of plant extract protein as determined by Western analysis (Table I). B.t.k. HD-1 concentrations for the plants containing the synthetic HD-1 gene averaged approximately 8200 ng (7200 and 9200 ng) of B.t.k. HD- 25 1 protein per mg of plant extract protein. This number compares well to the 5000 ng of HD-1 protein per mg of plant extract protein estimated by Western analysis. Likewise, plants containing the synthetic gene showed approximately a six-fold higher specific 30 activity than the corresponding plants containing the modified gene for these bioassays. In the Western

analysis the ratio was approximately 10 fold, again both are in good agreement. The level of *B.t.k.* protein in plants containing the wild-type HD-1 gene 5 (pMON9921) was too low to give a significant decrease in larval weight and hence was below a level that could be quantitated in this assay. In conclusion, the levels of *B.t.k.* HD-1 protein determined by both the bioassays and the Western analysis for these 10 plants containing the modified and synthetic genes agree, which demonstrates that the *B.t.k.* HD-1 protein produced by these plants is biologically active.

The levels of mRNA were determined in the plants containing the wild-type *B.t.k.* HD-1 gene (pMON9921) 15 and the modified gene (pMON5370) to establish whether the increased levels of protein production result from increased transcription or translation. mRNA from plants containing the synthetic gene could not be analyzed directly with the same DNA probe as used for the wild-type and modified genes because of the 20 numerous changes made in the coding sequence. mRNA was isolated and hybridized with a single-stranded DNA probe homologous to approximately the 5' 90 bp of the wild-type or modified gene coding sequences. The 25 hybrids were digested with S1 nuclease and the protected probe fragments analyzed by gel electrophoresis. Because the procedure used a large excess of probe and long hybridization time, the amount of protected probe is proportional to the amount of *B.t.k.* mRNA present in the sample. Two 30 plants expressing the modified gene (pMON5370) were

found to produce up to ten-fold more RNA than a plant expressing the wild-type gene (pMON9921).

5 The increased mRNA level from the modified gene is consistent with the result expected from the modifications introduced into this gene. However, this 10 fold increase in mRNA with the modified gene compared to the wild-type gene is in contrast to the 100 fold increase in *B.t.k.* protein from these genes  
10 in tobacco plants. If the two mRNAs were equally well translated then a 10 fold increase in stable mRNA would be expected to yield a 10 fold increase in protein. The higher increase in protein indicates that the modified gene mRNA is translated at about a  
15 10 fold higher efficiency than wild-type. Thus, about half of the total effect on gene expression can be explained by changes in mRNA levels and about half to changes in translational efficiency. This increase in translational efficiency is striking in that only  
20 about 9.5% of the codons have been changed in the modified gene; that is, this effect is clearly not due to wholesale codon usage changes. The increased translational efficiency could be due to changes in mRNA secondary structure that affect translation or to the removal of specific translational blockades due to  
25 specific codons that were changed.

The increased expression seen with the synthetic HD-1 gene was also seen with a synthetic HD-73 gene in tobacco. *B.t.k.* HD-73 was undetected in extracts of tobacco plants containing the wild-type truncated HD-30 73 gene (pMON5367), whereas *B.t.k.* HD-73 protein was easily detected in extracts from tobacco plants

5 containing the synthetic HD-73 gene of Figure 4 (pMON5383). Approximately 1000 ng of *B.t.k.* HD-73 protein was detected per mg of total soluble plant protein.

10 As described in Example 3 above, the *B.t.k.* HD-73 protein encoded in pMON5383 contains a small C-terminal extension of amino acids not encoded in the wild-type HD-73 protein. These extra amino acids had no effect on insect toxicity or on increased plant expression. A second synthetic HD-73 gene was constructed as described in Example 3 (Figure 8) and used to transform tobacco (pMON5390). Analysis of 15 plants containing pMON5390 showed that this gene was expressed at levels comparable to that of pMON5383 and that these plants had similar insecticidal efficacy.

20 In tobacco plants the synthetic HD-1 gene was expressed at approximately a 5-fold higher level than the synthetic HD-73 gene. However, this synthetic HD-73 gene still was expressed at least 100-fold better than the wild-type HD-73 gene. The HD-73 protein is approximately 5-fold more toxic to many insect pests than the HD-1 protein, so both synthetic HD-1 and HD-73 genes provide approximately comparable insecticidal efficacy in tobacco.

25 The full length *B.t.k.* HD-73 genes described in Example 3 were also incorporated into the plant transformation vector pMON893 so that they were expressed from the En 35S promoter. The synthetic/wild-type full length HD-73 gene of Figure 9 30 was incorporated into pMON893 to create pMON10505. The synthetic/modified full length HD-73 gene of

Figure 10 was incorporated into pMON893 to create pMON10526. The fully synthetic HD-73 gene of Figure 11 was incorporated into pMON893 to create pMON10518. 5 These vectors were used to obtain transformed tobacco plants, and the plants were analyzed for insecticidal efficacy and for *B.t.k.* HD-73 protein levels by Western blot or ELISA immunoassay.

Tobacco plants containing all three of these full 10 length *B.t.k.* genes produced detectable *B.t.k.* protein and showed 100% mortality of tobacco hornworm. This result is surprising in light of previous reported attempts to express the full length *B.t.k.* genes in transgenic plants. Vaeck et al. (1987) reported that 15 a full length *B.t.k. berliner* gene similar to our HD-1 gene could not be detectably expressed in tobacco. Barton et al. (1987) reported a similar result for another full length gene from *B.t.k.* HD-1 (the so called 4.5 kb gene), and further indicated that 20 tobacco callus containing this gene became necrotic, indicating that the full length gene product was toxic to plant cells. Fischhoff et al. (1987) reported that the full length *B.t.k.* HD-1 gene in tomato was poorly expressed compared to a truncated gene, and no plants that were fully toxic to tobacco hornworm could be 25 recovered. All three of the above reports indicated much higher expression levels and recovery of toxic plants if the respective *B.t.k.* genes were truncated. Adang et al. reported that the full length HD-73 gene yielded a few tobacco plants with some biological 30 activity (none were highly toxic) against hornworm and barely detectable *B.t.k.* protein. It was also noted

5 by them that the major *B.t.k.* mRNA in these plants was a truncated 1.7 kb species that would not encode a functional toxin. This indicated improper expression of the gene in tobacco. In contrast to all of these reports, the three full length *B.t.k.* HD-73 genes described above all lead to relatively high levels of protein and high levels of insect toxicity.

10 *B.t.k.* protein and mRNA levels in tobacco plants are shown in Table X for these three vectors. As can be seen from the table, the synthetic/wild-type gene (pMON10506) produces *B.t.k.* protein as about 0.01% of total soluble protein; the synthetic/modified gene produces *B.t.k.* as about 0.02% of total soluble protein; 15 and the fully synthetic gene produces *B.t.k.* as about 0.2% of total soluble protein. *B.t.k.* mRNA was analyzed in these plants by Northern blot analysis using the common 5' synthetic half of the genes as a probe. As shown in Table X, the increased protein levels can largely be attributed to increased mRNA 20 levels. Compared to the truncated modified and synthetic genes, this could indicate that the major contributors to increased translational efficiency are in the 5' half of the gene while the 3' half of the gene contains mostly determinants of mRNA stability. 25 The increased protein levels also indicate that increasing the amount of the full length gene that is synthetic or modified increases *B.t.k.* protein levels. Compared to the truncated synthetic *B.t.k.* HD-73 genes (pMON5383 or pMON5390), the fully synthetic gene 30 (pMON10518) produces as much or slightly more *B.t.k.* protein demonstrating that the full length genes are

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capable of being expressed at high levels in plants. These tobacco plants with high levels of full length 5 HD-73 protein show no evidence of abnormality and are fully fertile. The *B.t.k.* protein levels in these plants also produce the expected levels of insect toxicity based on feeding studies with beet armyworm or diet incorporation assays of plant extracts with tobacco budworm. The *B.t.k.* protein detected by 10 Western blot analysis in these tobacco plants often contains a varying amount of protein of about 80 kDa which is apparently a proteolytic fragment of the full length protein. The C-terminal half of the full length protein is known to be proteolytically 15 sensitive, and similar proteolytic fragments are seen from the full length gene in *E. coli* and *B.t.* itself. These fragments are fully insecticidal. The Northern analysis indicated that essentially all of the mRNA from these full length genes was of the expected full 20 length size. There is no evidence of truncated mRNAs that could give rise to the 80 kDa protein fragment. In addition, it is possible that the fragment is not present in intact plant cells and is merely due to proteolysis during extraction for immunoassay.

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Table X  
Full Length B.t.k. HD-73 Protein and  
mRNA Levels in Transgenic Tobacco Plants

5

	Gene description	Vector	B.t.k. protein concentration	Relative B.t.k. mRNA level
	Synthetic/wild type	pMON10506	>100	0.5
10	Synthetic/modified	pMON10526	400	1
	Fully synthetic	pMON10518	>2000	40

Thus, there is no serious impediment to producing high levels of *B.t.k.* HD-73 protein in plants from synthetic genes, and this is expected to be true of other full length lepidopteran active genes such as *B.t.k.* HD-1 or *B.t. entomocidus*. The fully synthetic *B.t.k.* HD-1 gene of Example 3 has been assembled in plant transformation vectors such as pMON893.

The fully synthetic gene in pMON10518 was also utilized in another plant vector and analyzed in tobacco plants. Although the CaMV35S promoter is generally a high level constitutive promoter in most plant tissues, the expression level of genes driven by the CaMV35S promoter is low in floral tissue relative to the levels seen in leaf tissue. Because the economically important targets damaged by some insects are the floral parts or derived from floral parts (e.g., cotton squares and bolls, tobacco buds, tomato buds and fruit), it may be advantageous to increase the expression of *B.t.* protein in these tissues over that obtained with the CaMV35S promoter.

The 35S promoter of Figwort Mosaic Virus (FMV) is analogous to the CaMV35S promoter. This promoter has been isolated and engineered into a plant transformation vector analogous to pMON893. Relative to the CaMV promoter, the FMV 35S promoter is highly expressed in the floral tissue, while still providing similar high levels of gene expression in other tissues such as leaf. A plant transformation vector, 5 pMON10517, was constructed in which the full length synthetic *B.t.k.* HD-73 gene of Figure 11 was driven by the FMV 35S promoter. This vector is identical to pMON10518 of Example 3 except that the FMV promoter is substituted for the CaMV promoter. Tobacco plants 10 transformed with pMON10517 and pMON10518 were obtained and compared for expression of the *B.t.k.* protein by Western blot or ELISA immunoassay in leaf and floral tissue. This analysis showed that pMON10517 containing the FMV promoter expressed the full length 15 HD-73 protein at higher levels in floral tissue than pMON10518 containing the CaMV promoter. Expression of the full length *B.t.k.* HD-73 protein from pMON10517 in leaf tissue is comparable to that seen with the most highly expressing plants containing pMON10518. 20 However, when floral tissue was analyzed, tobacco plants containing pMON10518 that had high levels of *B.t.k.* protein in leaf tissue did not have detectable *B.t.k.* protein in the flowers. On the other hand, flowers of tobacco plants containing pMON10517 had 25 levels of *B.t.k.* protein nearly as high as the levels in leaves at approximately 0.05% of total soluble 30 protein. This analysis showed that the FMV promoter

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could be used to produce relatively high levels of  
B.t.k. protein in floral tissue compared to the CaMV  
promoter.

5

b) Tomato.

The wild-type, modified and synthetic B.t.k. HD-1 genes tested in tobacco were introduced into other plants to demonstrate the broad utility of this 10 invention. Transgenic tomatoes were produced which contain these three genes. Data show that the increased expression observed with the modified and synthetic gene in tobacco also extends to tomato. Whereas the B.t.k. HD-1 protein is only barely 15 detectable in plants containing the wild type HD-1 gene (pMON9921), B.t.k. HD-1 was readily detected and the levels determined for plants containing the modified (pMON5370) or synthetic (pMON5377) genes. Expression levels for the plants containing the wild- 20 type, modified and synthetic HD-1 genes were approximately 10, 100 and 500 ng per mg of total plant extract (see Table XI below). The increase in B.t.k. HD-1 protein for the modified gene accounted for the majority of increase observed; 10 fold higher than the 25 plants containing the wild-type gene, compared to only an additional five-fold increase for plants containing the synthetic gene. Again the site-directed changes made in the modified gene are the major contributors to the increased expression of B.t.k. HD-1.

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Table XI

## B.t.k. HD-1 Expression in Transgenic Tomato Plants

Gene	B.t.k. Protein*	Fold Increase	
		in B.t.k.	
<u>Description</u>	<u>Vector</u>	<u>Concentration</u>	<u>Expression</u>
Wild type	pMON9921	10	1
Modified	pMON5370	100	10
Synthetic	pMON5377	500	50

15 \* B.t.k. HD-1 protein concentrations are expressed in ng/mg of total soluble plant protein. Data for plants containing the wild-type gene are estimates from mRNA levels and protein levels determined by ELISA.

20 These differences in B.t.k. HD-1 expression were confirmed with bioassays against tobacco hornworm and beet armyworm. Leaves from tomato plants containing each of these genes controlled tobacco hornworm damage and produced 100% mortality. With beet armyworm, leaves from plants containing the wild-type HD-1 gene (pMON9921) showed significant damage, leaves from plants containing the modified gene (pMON5370) showed less damage and leaves from plants containing the synthetic gene (pMON5377) were completely protected (see Table XII below).

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30

Table XIII

## Protection of Tomato Plants from 5 Tobacco Hornworm and Beet Armyworm

	Gene	<u>Description</u>	Tobacco Hornworm		Beet Armyworm	
			<u>Vector</u>	<u>Damage*</u>	<u>Vector</u>	<u>Damage*</u>
10	None	None		NL		NL
	Wild type	pMON9921		0		3
	Modified	pMON5370		0		1
	Synthetic	pMON5377		0		0

15 \* Damage was rated as shown in Table IX.

The generality of the synthetic gene approach was extended in tomato with a synthetic *B.t.k.* HD-73 gene.

20 In tomato, extracts from plants containing the wild-type truncated HD-73 gene (pMON5367) showed no detectable HD-73 protein. Extracts from plants containing the synthetic HD-73 gene (pMON5383) showed high levels of *B.t.k.* HD-73 protein, approximately 2000 ng per mg of plant extract protein. These data clearly demonstrate that the changes made in the 25 synthetic HD-73 gene lead to dramatic increases in the expression of the HD-73 protein in tomato as well as in tobacco

In contrast to tobacco, the synthetic HD-73 gene in tomato is expressed at approximately 4-fold to 5-fold higher levels than the synthetic HD-1 gene. Because the HD-73 protein is about 5-fold more active than the

5 HD-1 protein against many insect pests including Heliothis species, the increased expression of synthetic HD-73 compared to synthetic HD-1 corresponds to about a 25-fold increased insecticidal efficacy in tomato.

10 In order to determine the mechanisms involved in the increased expression of modified and synthetic B.t.k. HD-1 genes in tomato, S1 nuclease analysis of mRNA levels from transformed tomato plants was performed. As indicated above, a similar analysis had been performed with tobacco plants, and this analysis showed that the modified gene produced up to 10-fold more mRNA than the wild-type gene. The analysis in 15 tomato utilized a different DNA probe that allowed the analysis of wild-type (pMON9921), modified (pMON5370) and synthetic (pMON5377) HD-1 genes with the same probe. This probe was derived from the 5' untranslated region of the CaMV35S promoter in pMON893 20 that was common to all three of these vectors (pMON9921, pMON5370 and pMON5377). This S1 analysis indicated that B.t.k. mRNA levels from the modified gene were 3 to 5 fold higher than for the wild-type gene, and that mRNA levels for the synthetic gene were about 2 to 3 fold higher than for the modified gene. 25 Three independent transformants were analyzed for each gene. Compared to the fold increases in B.t.k. HD-1 protein from these genes in tomato shown in Table XI, these mRNA increases can explain about half of the total protein increase as was seen in tobacco for the 30 wild-type and modified genes. For tomato the total mRNA increase from wild-type to synthetic is about 6

to 15 fold compared to a protein increase of about 50 fold. This result is similar to that seen for tobacco in comparing the wild-type and modified genes, and it 5 extends to the synthetic gene as well. That is, about half of the total fold increase in *B.t.k.* protein from wild-type to modified genes can be explained by mRNA increases and about half to enhanced translational efficiency. The same is also true in comparing the 10 modified gene to the synthetic gene. Although there is an additional increase in RNA levels, this mRNA increase can explain only about half of the total protein increase.

The full length *B.t.k.* genes described above were 15 also used to transform tomato plants and these plants were analyzed for *B.t.k.* protein and insecticidal efficacy. The results of this analysis are shown in Table XIII. Plants containing the synthetic/wild-type gene (pMON10506) produce the *B.t.k.* HD-73 protein at 20 levels of about 0.01% of their total soluble protein. Plants containing the synthetic/modified gene (pMON10526) produce about 0.04% *B.t.k.* protein, and plants containing the fully synthetic gene (pMON10518) produce about 0.2% *B.t.k.* protein. These results are very similar to the tobacco plant results for the same 25 genes. mRNA levels estimated by Northern blot analysis in tomato also increase in parallel with the protein level increase. As for tobacco with these three genes, most of the protein increase can be attributed to increased mRNA with a small component of 30 translational efficiency increase indicated for the fully synthetic gene. The highest levels of full

length *B.t.k.* protein (from pMON10518) are comparable to or just slightly lower than the highest levels observed for the truncated HD-73 genes (pMON5383 and pMON5390). Tomato plants expressing these full length genes have the insecticidal activity expected for the observed protein levels as determined by feeding assays with beet armyworm or by diet incorporation of plant extracts with tobacco hornworm.

10

Table XIII

Full Length *B.t.k.* HD-73 Protein and mRNA Levels in Transgenic Tomato Plants

15

	Gene description	Vector	<i>B.t.k.</i> protein concentration	Relative <i>B.t.k.</i> mRNA level
	Synthetic/wild type	pMON10506	100	1
20	Synthetic/modified	pMON10526	400	2-4
	Fully synthetic	pMON10518	2000	10

c) Cotton.

The generality of the increased expression of *B.t.k.* HD-1 and *B.t.k.* HD-73 by use of the modified and synthetic genes was extended to cotton. Transgenic calli were produced which contain the wild type (pMON9921) and the synthetic HD-1 (pMON5377) genes. Here again the *B.t.k.* HD-1 protein produced from calli containing the wild-type gene was not detected, whereas calli containing the synthetic HD-1

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gene expressed the HD-1 protein at easily detectable levels. The HD-1 protein was produced at approximately 1000 ng/mg of plant calli extract protein. Again, to ensure that the protein produced by the transgenic cotton calli was biologically active and that the increased expression observed with the synthetic gene translated to increased biological activity, extracts of cotton calli were made in a similar manner as described for tobacco plants, except that the calli was first dried between Whatman filter paper to remove as much of the water as possible. The dried calli were then ground in liquid nitrogen and ground in 100 mM sodium carbonate buffer, pH 10. Approximately 0.5 ml aliquotes of this material was applied to tomato leaves with a paint brush. After the leaf dried, five tobacco hornworm larvae were applied to each of two leaf samples. Leaves painted with extract from control calli were completely destroyed. Leaves painted with extract from calli containing the wild-type HD-1 gene (pMON9921) showed severe damage. Leaves painted with extract from calli containing the synthetic HD-1 gene (pMON5377) showed no damage (see Table XIV below).

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Table XIV

5      Protection against Tobacco Hornworm by Tomato Leaves  
Painted with Extracts Prepared from Cotton Calli  
Containing a Control, the Wild-Type B.t.k. HD-1 Gene,  
Synthetic HD-1 Gene or Synthetic HD-73 Gene

	Gene		Tobacco Hornworm
10	<u>Description</u>	<u>Vector</u>	<u>Damage*</u>
	Control	Control	NL
	Wild type HD-1	pMON9921	3
	Synthetic HD-1	pMON5377	0
15	Synthetic HD-73	pMON5383	0

\* Damage was rated as shown in Table VIII.

20      Cotton calli were also produced containing another synthetic gene, a gene encoding B.t.k. HD-73. The preparation of this gene is described in Example 3. Calli containing the synthetic HD-73 gene produced the corresponding HD-73 protein at even higher levels than the calli which contained the synthetic HD-1 gene. Extracts made from calli containing the HD-73

25      synthetic gene (pMON5383) showed complete control of tobacco hornworm when painted onto tomato leaves as described above for extracts containing the HD-1 protein. (See Table XIV).

30      Transgenic cotton plants containing the synthetic B.t.k. HD-1 gene (pMON5377) or the synthetic B.t.k. HD-73 gene (pMON5383) have also been examined. These

5 plants produce the HD-1 or HD-73 proteins at levels comparable to that seen in cotton callus with the same genes and comparable to tomato and tobacco plants with these genes. For either synthetic truncated HD-1 or HD-73 genes, cotton plants expressing *B.t.k.* protein at 1000 to 2000 ng/mg total protein (0.1% to 0.2%) were recovered at a high frequency. Insect feeding assays were performed with leaves from cotton plants 10 expressing the synthetic HD-1 or HD-73 genes. These leaves showed no damage (rating of 0) when challenged with larvae of cabbage looper (*Trichoplusia ni*), and only slight damage when challenged with larvae of beet armyworm (*Spodoptera exigua*). Damage ratings are as 15 defined in Table VIII above. This demonstrated that cotton plants as well as calli expressed the synthetic HD-1 or HD-73 genes at high levels and that those plants were protected from damage by Lepidopteran insect larvae.

20 Transgenic cotton plants containing either the synthetic truncated HD-1 gene (pMON5377) or the synthetic truncated HD-73 gene (pMON5383) were also assessed for protection against cotton bollworm at the whole plant level in the greenhouse. This is a more realistic test of the ability of these plants to 25 produce an agriculturally acceptable level of control. The cotton bollworm (*Heliothis zea*) is a major pest of cotton that produces economic damage by destroying terminals, squares and bolls, and protection of these fruiting bodies as well as the leaf tissue will be 30 important for effective insect control and adequate crop protection. To test the protection afforded to

whole plants, R1 progeny of cotton plants expressing high levels of either *B.t.k.* HD-1 (pMON5377) or *B.t.k.* HD-73 (pMON5383) were assayed by applying 10-15 eggs of cotton bollworm per boll or square to the 20 uppermost squares or bolls on each plant. At least 12 plants were analyzed per treatment. The hatch rate of the eggs was approximately 70%. This corresponds to very high insect pressure compared to numbers of larvae per plant seen under typical field conditions. Under these conditions 100% of the bolls on control cotton plants were destroyed by insect damage. For the transgenics, significant boll protection was observed. Plants containing pMON5377 (HD-1) had 70-75% of the bolls survive the intense pressure of this assay. Plants containing pMON5383 (HD-73) had 80% to 90% boll protection. This is likely to be a consequence of the higher activity of HD-73 protein against cotton bollworm compared to HD-1 protein. In cases where the transgenic plants were damaged by the insects, the surviving larvae were delayed in their development by at least one instar.

Therefore, the increased expression obtained with the modified and synthetic genes is not limited to any one crop; tobacco, tomato and cotton calli and cotton plants all showed drastic increases in *B.t.k.* expression when the plants/callus were produced containing the modified or synthetic genes. Likewise, the utility of changes made to produce the modified and synthetic *B.t.k.* HD-1 gene is not limited to the HD-1 gene. The synthetic HD-73 gene in all three species also showed drastic increases in expression.

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In summary, it has been demonstrated that: (1) the genetic changes made in the HD-1 modified gene lead to very significant increases in B.t.k. HD-1 expression;

5 further five-fold increase in B.t.k. HD-1 expression; (3) the changes incorporated into the modified HD-1 gene accounted for the majority of the increased B.t.k. expression observed with the synthetic gene; 10 (4) the increased expression was demonstrated in three different plants -- tobacco plants, tomato plants and cotton calli and cotton plants; (5) the increased expression as observed by Western analysis also correlated with similar increases in bioactivity, 15 showing that the B.t.k. HD-1 proteins produced were comparably active; (6) when the method of the present invention used to design the synthetic HD-1 gene was employed to design a synthetic HD-73 gene it also was expressed at much higher levels in tobacco, tomato and 20 cotton than the wild-type equivalent gene with consequent increases in bioactivity; (7) a fully synthetic full length B.t.k. gene was expressed at levels comparable to synthetic truncated genes.

25 Example 5 -- Synthetic B.t. tenebrionis Gene in  
Tobacco, Tomato and Potato

Referring to Figure 12, a synthetic gene encoding a Coleopteran active toxin is prepared by making the indicated changes in the wild-type gene of B.t. 30 tenebrionis or de novo synthesis of the synthetic

5 an intermediate plant transformation vector such as pMON893: Plasmid pMON893 containing the synthetic *B.t.t.* gene is then inserted into a suitable disarmed *Agrobacterium* strain such as *A. tumefaciens* ACO.

Transformation and Regeneration of Potato

10 Sterile shoot cultures of Russet Burbank are maintained in vials containing 10 ml of PM medium (Murashige and Skoog (MS) inorganic salts, 30 g/l sucrose, 0.17 g/l  $\text{NaH}_2\text{PO}_4\text{H}_2\text{O}$ , 0.4 mg/l thiamine-HCl, and 100 mg/l myo-inositol, solidified with 1 g/l Gelrite at pH 6.0). When shoots reached approximately 15 5 cm in length, stem internode segments of 7-10 mm are excised and smeared at the cut ends with a disarmed *Agrobacterium tumefaciens* vector containing the synthetic *B.t.t.* gene from a four day old plate culture. The stem explants are co-cultured for three 20 days at 23°C on a sterile filter paper placed over 1.5 ml of a tobacco cell feeder layer overlaid on 1/10 P medium (1/10 strength MS inorganic salts and organic addenda without casein as in Jarret et al. (1980), 30 g/l sucrose and 8.0 g/l agar). Following co-culture 25 the explants are transferred to full strength P-1 medium for callus induction, composed of MS inorganic salts, organic additions as in Jarret et al. (1980) with the exception of casein, 3.0 mg/l benzyladenine (BA), and 0.01 mg/l naphthaleneacetic acid (NAA) (Jarret, et al., 1980). Carbenicillin (500 mg/l) is 30 included to inhibit bacterial growth, and 100 mg/l

kanamycin is added to select for transformed cells. After four weeks the explants are transferred to medium of the same composition but with 0.3 mg/l gibberellic acid (GA3) replacing the BA and NAA (Jarret et al., 1981) to promote shoot formation. Shoots begin to develop approximately two weeks after transfer to shoot induction medium; these are excised and transferred to vials of PM medium for rooting.

5 Shoots are tested for kanamycin resistance conferred by the enzyme neomycin phosphotransferase II, by placing a section of the stem onto callus induction medium containing MS organic and inorganic salts, 30 g/l sucrose, 2.25 mg/l BA, 0.186 mg/l NAA, 10 mg/l

10 GA3 (Webb, et al., 1983) and 200 mg/l kanamycin to

15 select for transformed cells.

The synthetic *B.t.t.* gene described in figure 12, was placed into a plant expression vector as described in example 5. The plasmid has the following characteristics; a synthetic BglII fragment having approximately 1800 base pairs was inserted into pMON893 in such a manner that the enhanced 35S promoter would express the *B.t.t.* gene. This construct, pMON1982, was used to transform both tobacco and tomato. Tobacco plants, selected as

20 kanamycin resistant plants were screened with rabbit anti-*B.t.t.* antibody. Cross-reactive material was detected at levels predicted to be suitable to cause mortality to CPB. These target insects will not feed on tobacco, but the transgenic tobacco plants do

25

30 demonstrate that the synthetic gene does improve expression of this protein to detectable levels.

5 Tomato plants with the pMON1982 construct were determined to produce *B.t.t.* protein at levels insecticidal to CPB. In initial studies, the leaves of four plants (5190, 5225, 5328 and 5133) showed little or no damage when exposed to CPB larvae (damage rating of 0-1 on a scale of 0 to 4 with 4 as no leaf remaining). Under these conditions the control leaves were completely eaten. Immunological analysis of 10 these plants confirmed the presence of material cross-reactive with anti-*B.t.t.* antibody. Levels of protein expression in these plants were estimated at approximately 1 to 5 ng of *B.t.t.* protein in 50 ug of total extractable protein. A total of 17 tomato 15 plants (17 of 65 tested) have been identified which demonstrate protection of leaf tissue from CPB (rating of 0 or 1) and show good insect mortality.

20 Results similar to those seen in tobacco and tomato with pMON1982 were seen with pMON1984 in the same plant species. pMON1984 is identical to pMON1982 except that the synthetic protease inhibitor (CMTI) is fused upstream of the native proteolytic cleavage site. Levels of expression in tobacco were estimated to be similar to pMON1982, between 10-15 ng per 50ug 25 of total soluble protein.

25 Tomato plants expressing pMON1984 have been identified which protect the leaves from ingestion by CPB. The damage rating was 0 with 100% insect mortality.

30 Potato was transformed as described in example 5 with a vector similar to pMON1982 containing the enhanced CaMV35S/synthetic *B.t.t.* gene. Leaves of

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5 potato plants transformed with this vector, were screened by CPB insect bioassay. Of the 35 plants tested, leaves from 4 plants, 16a, 13c, 13d, and 23a were totally protected when challenged. Insect bioassays with leaves from three other plants, 13e, 1a, and 13b, recorded damage levels of 1 on a scale of 0 to 4 with 4 being total devestation of the leaf material. Immunological analysis confirmed the 10 presence of *B.t.t.* cross-reactive material in the leaf tissue. The level of *B.t.t.* protein in leaf tissue of plant 16a (damage rating of 0) was estimated at 20-50 ng of *B.t.t.* protein/50 ug of total soluble protein. The levels of *B.t.t.* protein seen in 16a tissue was 15 consistent with its biological activity. Immunological analysis of 13e and 13b (tissue which scored 1 in damage rating) reveal less protein (5-10 ng/50 ug of total soluble protein) than in plant 16a. Cuttings of plant 16a were challenged with 50 to 200 20 eggs of CPB in a whole plant assay. Under these conditions 16a showed no damage and 100% mortality of insects while control potato plants were heavily damaged.

25 Example 6 -- Synthetic *B.t.k.* P2 Protein Gene

30 The P2 protein is a distinct insecticidal protein produced by some strains of *B.t.* including *B.t.k.* HD-1. It is characterized by its activity against both lepidopteran and dipteran insects (Yamamoto and Iizuka, 1983). Genes encoding the P2 protein have been isolated and characterized (Donovan et al.,

1988). The P2 proteins encoded by these genes are approximately 600 amino acids in length. These 5 proteins share only limited homology with the lepidopteran specific P1 type proteins, such as the *B.t.k.* HD-1 and HD-73 proteins described in previous examples.

The P2 proteins have substantial activity against a variety of lepidopteran larvae including cabbage 10 looper, tobacco hornworm and tobacco budworm. Because they are active against agronomically important insect pests, the P2 proteins are a desirable candidate in the production of insect tolerant transgenic plants either alone or in combination with the other *B.t.* 15 toxins described in the above examples. In some plants, expression of the P2 protein alone might be sufficient to provide protection against damaging insects. In addition, the P2 proteins might provide protection against agronomically important dipteran pests. In other cases, expression of P2 together with 20 the *B.t.k.* HD-1 or HD-73 protein might be preferred. The P2 proteins should provide at least an additive level of insecticidal activity when combined with the crystal protein toxin of *B.t.k.* HD-1 or HD-73, and the 25 combination may even provide a synergistic activity. Although the mode of action of the P2 protein is unknown, its distinct amino acid sequence suggests that it functions differently from the *B.t.k.* HD-1 and HD-73 type of proteins. Production of two insect tolerance proteins with different modes of action in 30 the same plant would minimize the potential for development of insect resistance to *B.t.* proteins in

plants. The lack of substantial DNA homology between P2 genes and the HD-1 and HD-73 genes minimizes the potential for recombination between multiple insect tolerance genes in the plant chromosome.

The genes encoding the P2 protein although distinct in sequence from the *B.t.k.* HD-1 and HD-73 genes share many common features with these genes. In particular, the P2 protein genes have a high A+T content (65%), multiple potential polyadenylation signal sequences (26) and numerous ATTAA sequences (10). Because of its overall similarity to the poorly expressed wild-type *B.t.k.* HD-1 and HD-73 genes, the same problems are expected in expression of the wild-type P2 gene as were encountered with the previous examples. Based on the above-described method for designing the synthetic *B.t.* genes, a synthetic P2 gene has been designed which gene should be expressed at adequate levels for protection in plants. A comparision of the wild-type and synthetic P2 genes is shown in Figure 13.

Example 7 -- Synthetic *B.t.* *Entomocidus* Gene

The *B.t. entomocidus* ("Btent") protein is a distinct insecticidal protein produced by some strains of *B.t.* bacteria. It is characterized by its high level of activity against some lepidopterans that are relatively insensitive to *B.t.k.* HD-1 and HD-73 such as *Spodoptera* species including beet armyworm (Visser et al., 1988). Genes encoding the Btent protein have been isolated and characterized (Honee et al, 1988). The Btent proteins encoded by these genes are

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approximately the same length as *B.t.k.* HD-1 and HD-73. These proteins share only 68% amino acid homology with the *B.t.k.* HD-1 and HD-73 proteins. It is likely that only the N-terminal half of the Btent protein is required for insecticidal activity as is the case for HD-1 and HD-73. Over the first 625 amino acids, Btent shares only 38% amino acid homology with HD-1 and HD-73.

Because of their higher activity against *Spodoptera* species that are relatively insensitive to HD-1 and HD-73, the Btent proteins are a desirable candidate for the production of insect tolerant transgenic plants either alone or in combination with the other *B.t.* toxins described in the above examples. In some plants production of Btent alone might be sufficient to control the agronomically important pests. In other plants, the production of two distinct insect tolerance proteins would provide protection against a wider array of insects. Against those insects where both proteins are active, the combination of the *B.t.k.* HD-1 or HD-73 type protein plus the Btent protein should provide at least additive insecticidal efficacy, and may even provide a synergistic activity. In addition, because of its distinct amino acid sequence, the Btent protein may have a different mode of action than HD-1 or HD-73. Production of two insecticidal proteins in the same plant with different modes of action would minimize the potential for development of insect resistance to *B.t.* proteins in plants. The relative lack of DNA sequence homology with the *B.t.k.* type genes minimizes the potential for

recombination between multiple insect tolerance genes

5       The genes encoding the Btent protein although  
distinct in sequence from the B.t.k. HD-1 and HD-73  
genes share many common features with these genes. In  
particular, the Btent protein genes have a high A+T  
content (62%), multiple potential polyadenylation  
signal sequences (39 in the full length coding  
10      sequence and 27 in the first 1875 nucleotides that is  
likely to encode the active toxic fragment) and  
numerous ATTAA sequences (16 in the full length coding  
sequence and 12 in the first 1875 nucleotides).  
Because of its overall similarity to the poorly  
15      expressed wild type B.t.k. HD-1 and HD-73 genes, the  
wild-type Btent genes are expected to exhibit similar  
problems in expression as were encountered with the  
wild-type HD-1 and HD-73 genes. Based on the above-  
described method used for designing the other  
20      synthetic B.t. genes, a synthetic Btent gene has been  
designed which gene should be expressed at adequate  
levels for protection in plants. A comparision of the  
wild type and synthetic Btent genes is shown in  
Figure 14.

25      Example 8 -- Synthetic B.t.k. Genes for Expression  
in Corn

30      High level expression of heterologous genes in corn  
cells has been shown to be enhanced by the presence of  
a corn gene intron (Callis et al., 1987). Typically  
these introns have been located in the 5' untranslated

region of the chimeric gene. It has been shown that the CaMV35S promoter and the NOS 3' end function efficiently in the expression of heterologous genes in 5 corn cells (Fromm et al., 1986).

Referring to Figure 15, a plant expression cassette vector (pMON744) was constructed that contains these sequences. Specifically the expression cassette contains the enhanced CaMV 35S promoter followed by 10 intron 1 of the corn Adh1 gene (Callis et al., 1987). This is followed by a multilinker cloning site for insertion of coding sequences; this multilinker contains a BglII site among others. Following the multilinker is the NOS 3' end. pMON744 also contains 15 the selectable marker gene 35S/NPTII/NOS 3' for kanamycin selection of transgenic corn cells. In addition, pMON744 has an *E. coli* origin of replication and an ampicillin resistance gene for selection of the plasmid in *E. coli*.

20 Five *B.t.k.* coding sequences described in the previous examples were inserted into the BglII site of pMON744 for corn cell expression of *B.t.k.* The coding sequences inserted and resulting vectors were:

- 25 1. Wild type *B.t.k.* HD-1 from pMON9921 to make pMON8652.
2. Modified *B.t.k.* HD-1 from pMON5370 to make pMON8642.
3. Synthetic *B.t.k.* HD-1 from pMON5377 to make pMON8643.
- 30 4. Synthetic *B.t.k.* HD-73 from pMON5390 to make pMON8644.

5. Synthetic full length *B.t.k.* HD-73 from pMON10518 to make pMON10902.

5 pMON8652 (wild-type *B.t.k.* HD-1) was used to transform corn cell protoplasts and stably transformed kanamycin resistant callus was isolated. *B.t.k.* mRNA in the corn cells was analyzed by nuclease S1 protection and found to be present at a level  
10 comparable to that seen with the same wild-type coding sequence (pMON9921) in transgenic tomato plants.

15 pMON8652 and pMON8642 (modified HD-1) were used to transform corn cell protoplasts in a transient expression system. The level of *B.t.k.* mRNA was analyzed by nuclease S1 protection. The modified HD-1 gave rise to a several fold increase in *B.t.k.* mRNA compared to the wild-type coding sequence in the transiently transformed corn cells. This indicated that the modifications introduced into the *B.t.k.* HD-1  
20 gene are capable of enhancing *B.t.k.* expression in monocot cells as was demonstrated for dicot plants and cells.

25 pMON8642 (modified HD-1) and pMON8643 (synthetic HD-1) were used to transform Black Mexican Sweet (BMS) corn cell protoplasts by PEG-mediated DNA uptake, and stably transformed corn callus was selected by growth on kanamycin containing plant growth medium. Individual callus colonies that were derived from single transformed cells were isolated and propagated separately on kanamycin containing medium.

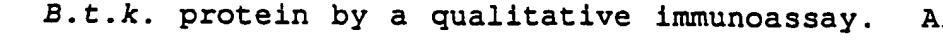
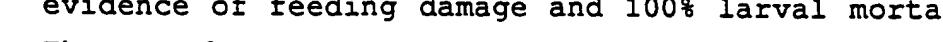
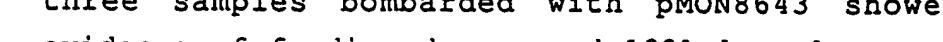
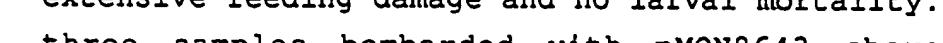
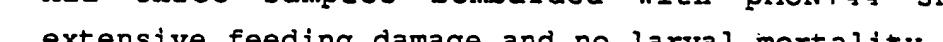
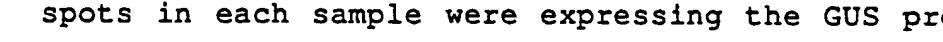
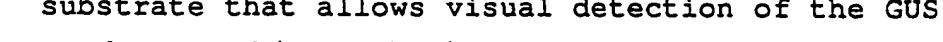
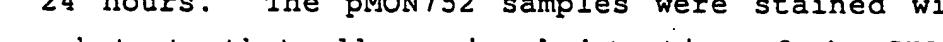
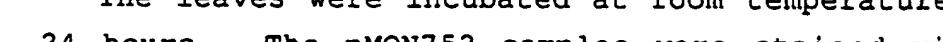
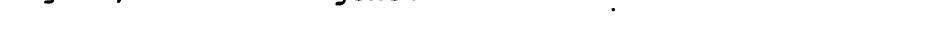
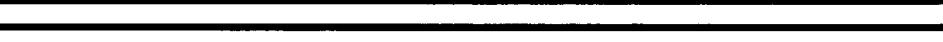
30 To assess the expression of the *B.t.k.* genes in these cells, callus samples were tested for insect

toxicity by bioassay against tobacco hornworm larvae. For each vector, 96 callus lines were tested by 5 bioassay. Portions of each callus were placed on sterile water agar plates, and five neonate tobacco hornworm larvae were added and allowed to feed for 4 days. For pMON8643, 100% of the larvae died after feeding on 15 of the 96 calli and these calli showed little feeding damage. For pMON8642, only 1 of the 96 10 calli was toxic to the larvae. This showed that the *B.t.k.* gene was being expressed in these samples at insecticidal levels. The observation that significantly more calli containing pMON8643 were toxic than for pMON8642 showed that significantly 15 higher levels of expression were obtained when the synthetic HD-1 coding sequence was contained in corn cells than when the modified HD-1 coding sequence was used, similar to the previous examples with dicot plants. A semiquantitative immunoassay showed that 20 the pMON8643 toxic samples had significantly higher *B.t.k.* protein levels than the pMON8642 toxic sample.

The 16 callus samples that were toxic to tobacco hornworm were also tested for activity against European corn borer. European corn borer is 25 approximately 40-fold less sensitive to the HD-1 gene product than is tobacco hornworm. Larvae of European corn borer were applied to the callus samples and allowed to feed for 4 days. Two of the 16 calli tested, both of which contained pMON8643 (synthetic HD- 30 1), were toxic to European corn borer larvae.

To assess the expression of the *B.t.k.* genes in differentiated corn tissue, another method of DNA delivery was used. Young leaves were excised from 5 corn plants, and DNA samples were delivered into the leaf tissue by microprojectile bombardment. In this system, the DNA on the microprojectiles is transiently expressed in the leaf cells after bombardment. Three DNA samples were used, and each DNA was tested in 10 triplicate.

1. pMON744, the corn expression vector with no



15 3. pMON752, a corn expression vector for the GUS gene, no *B.t.k.* gene.

The leaves were incubated at room temperature for 24 hours. The pMON752 samples were stained with a 20 substrate that allows visual detection of the GUS gene product. This analysis showed that over one hundred spots in each sample were expressing the GUS product and the triplicate samples showed very similar levels of GUS expression. For the pMON744 and pMON8643 samples 5 larvae of tobacco hornworm were 25 added to each leaf and allowed to feed for 48 hours. All three samples bombarded with pMON744 showed extensive feeding damage and no larval mortality. All three samples bombarded with pMON8643 showed no evidence of feeding damage and 100% larval mortality. 30 The samples were also assayed for the presence of *B.t.k.* protein by a qualitative immunoassay. All of

the pMON8643 samples had detectable *B.t.k.* protein. These results demonstrated that the the synthetic *B.t.k.* gene was expressed in differentiated corn plant tissue at insecticidal levels.

Example 9 -- Synthetic Potato Leaf Roll Virus Coat Protein Gene

10 Expression in plants of the coat protein genes from a variety of plant viruses has proven to be an effective method of engineering resistance to these viruses. In order to achieve virus resistance, it is important to express the viral coat protein at an 15 effective level. For many plant virus coat protein genes, this has not proved to be a problem. However, for the coat protein gene from potato leaf roll virus (PLRV), expression of the coat protein has been observed to be low relative to other coat protein 20 genes, and this lower level of protein has not led to optimal resistance to PLRV.

The gene for PLRV coat protein is shown in Figure 16. Referring to Figure 16, the upper line of sequence shows the gene as it was originally engineered for plant expression in vector pMON893. 25 The gene was contained on a 749 nucleotide *Bgl*II-*Eco*RI fragment with the coding sequence contained between nucleotides 20 and 643. This fragment also contained 19 nucleotides of 5' noncoding sequence and 104 nucleotides of 3' noncoding sequence. This PLRV coat 30 protein gene was relatively poorly expressed in plants compared to other viral coat protein genes.

A synthetic gene was designed to improve plant expression of the PLRV coat protein. Referring again to Figure 16, the changes made in the synthetic PLRV gene are shown in the lower line. This gene was designed to encode exactly the same protein as the naturally occurring gene. Note that the beginning of the synthetic gene is at nucleotide 14 and the end of the sequence is at nucleotide 654. The coding sequence for the synthetic gene is from nucleotide 20 to 643 of the figure. The changes indicated just upstream and downstream of these endpoints serve only to introduce convenient restriction sites just outside the coding sequence. Thus the size of the synthetic gene is 641 nucleotides which is smaller than the naturally occurring gene. The synthetic gene is smaller because substantially all of the noncoding sequence at both the 5' and 3' ends, except for segments encoding the BglII and EcoRI restriction sites has been removed.

The synthetic gene differs from the naturally occurring gene in two main respects. First, 41 individual codons within the coding sequence have been changed to remove nearly all codons for a given amino acid that constitute less than about 15% of the codons for that amino acid in a survey of dicot plant genes. Second, the 5' and 3' noncoding sequences of the original gene have been removed. Although not strictly conforming to the algorithm described in Figure 1, a few of the codon changes and especially the removal of the long 3' noncoding region is consistent with this algorithm.

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The original PLRV sequence contains two potential plant polyadenylation signals (AACCAA and AAGCATT) and both of these occur in the 3' noncoding sequence. 5 that has been removed in the synthetic gene. The original PLRV gene also contains one ATTAA sequence. This is also contained in the 3' noncoding sequence, and is in the midst of the longest stretch of uninterrupted A+T in the gene (a stretch of 7 A+T 10 nucleotides). This sequence was removed in the synthetic gene. Thus, sequences that the algorithm of Figure 1 targets for change have been changed in the synthetic PLRV coat protein gene by removal of the 3' noncoding segment. Within the coding sequence, codon 15 changes were also made to remove three other regions of sequence described above. In particular, two regions of 5 consecutive A+T and one region of 5 consecutive G+C within the coding sequence have been removed in the synthetic gene.

20 The synthetic PLRV coat protein gene is cloned in a plant transformation vector such as pMON893 and used to transform potato plants as described above. These plants express the PLRV coat protein at higher levels than achieved with the naturally occurring gene, and 25 these plants exhibit increased resistance to infection by PLRV.

Example 10 -- Expression of Synthetic B.t. Genes  
with RUBISCO Small Subunit Promoters and  
Chloroplast Transit Peptides

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The genes in plants encoding the small subunit of RUBISCO (SSU) are often highly expressed, light regulated and sometimes show tissue specificity. These expression properties are largely due to the 10 promoter sequences of these genes. It has been possible to use SSU promoters to express heterologous genes in transformed plants. Typically a plant will contain multiple SSU genes, and the expression levels and tissue specificity of different SSU genes will be 15 different. The SSU proteins are encoded in the nucleus and synthesized in the cytoplasm as precursors that contain an N-terminal extension known as the chloroplast transit peptide (CTP). The CTP directs the precursor to the chloroplast and promotes the 20 uptake of the SSU protein into the chloroplast. In this process, the CTP is cleaved from the SSU protein. These CTP sequences have been used to direct heterologous proteins into chloroplasts of transformed plants.

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The SSU promoters might have several advantages for expression of B.t.k. genes in plants. Some SSU promoters are very highly expressed and could give rise to expression levels as high or higher than those observed with the CaMV35S promoter. The tissue distribution of expression from SSU promoters is 30 different from that of the CaMV35S promoter, so for control of some insect pests, it may be advantageous

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to direct the expression of *B.t.k.* to those cells in which SSU is most highly expressed. For example, 5 although relatively constitutive, in the leaf the CaMV35S promoter is more highly expressed in vascular tissue than in some other parts of the leaf, while most SSU promoters are most highly expressed in the mesophyll cells of the leaf. Some SSU promoters also are more highly tissue specific, so it could be 10 possible to utilize a specific SSU promoter to express *B.t.k.* in only a subset of plant tissues, if for example *B.t.* expression in certain cells was found to be deleterious to those cells. For example, for control of Colorado potato beetle in potato, it may be 15 advantageous to use SSU promoters to direct *B.t.t.* expression to the leaves but not to the edible tubers.

Utilizing SSU CTP sequences to localize *B.t.* proteins to the chloroplast might also be 20 advantageous. Localization of the *B.t.* to the chloroplast could protect the protein from proteases found in the cytoplasm. This could stabilize the *B.t.* protein and lead to higher levels of accumulation of active protein. *B.t.* genes containing the CTP could 25 be used in combination with the SSU promoter or with other promoters such as CaMV35S.

A variety of plant transformation vectors were constructed for the expression of *B.t.k.* genes utilizing SSU promoters and SSU CTPs. The promoters and CTPs utilized were from the petunia SSU11a gene 30 described by Turner et al. (1986) and from the *Arabidopsis ats1A* gene (an SSU gene) described by

Krebbers et al. (1988) and by Elionor et al. (1989). The petunia SSU11a promoter was contained on a DNA fragment that extended approximately 800 bp upstream of the SSU coding sequence. The *Arabidopsis* ats1A promoter was contained on a DNA fragment that extended approximately 1.8 kb upstream of the SSU coding sequence. At the upstream end convenient sites from the multilinker of pUC18 were used to move these promoters into plant transformation vectors such as pMON893. These promoter fragments extended to the start of the SSU coding sequence at which point an NcoI restriction site was engineered to allow insertion of the *B.t.* coding sequence, replacing the SSU coding sequence.

When SSU promoters were used in combination with their CTP, the DNA fragments extended through the coding sequence of the CTP and a small portion of the mature SSU coding sequence at which point an NcoI restriction site was engineered by standard techniques to allow the in frame fusion of *B.t.* coding sequences with the CTP. In particular, for the petunia SSU11a CTP, *B.t.* coding sequences were fused to the SSU sequence after amino acid 8 of the mature SSU sequence at which point the NcoI site was placed. The 8 amino acids of mature SSU sequence were included because preliminary in vitro chloroplast uptake experiments indicated that uptake was of *B.t.* was observed only if this segment of mature SSU was included. For the *Arabidopsis* ats1A CTP, the complete CTP was included plus 24 amino acids of mature SSU sequence plus the sequence gly-gly-arg-val-asn-cys-met-gln-ala-met,

5 terminating in an NcoI site for *B.t.* fusion. This short sequence reiterates the native SSU CTP cleavage site (between the cys and met) plus a short segment surrounding the cleavage site. This sequence was included in order to insure proper uptake into chloroplasts. *B.t.* coding sequences were fused to this ats1A CTP after the met codon. In vitro uptake experiments with this CTP construction and other (non-  
10 *B.t.*) coding sequences showed that this CTP did target proteins to the chloroplast.

When CTPs were used in combination with the CaMV 35S promoter, the same CTP segments were used. They were excised just upstream of the ATG start sites of  
15 the CTP by engineering of BglII sites, and placed downstream of the CaMV35S promoter in pMON893, as BglII to NcoI fragments. *B.t.* coding sequences were fused as described above.

20 The wild type *B.t.k.* HD-1 coding sequence of pMON9921 (see Figure 1) was fused to the ats1A promoter to make pMON1925 or the ats1A promoter plus CTP to make pMON1921. These vectors were used to transform tobacco plants, and the plants were screened for activity against tobacco hornworm. No toxic plants were recovered. This is surprising in light of  
25 the fact that toxic plants could be recovered, albeit at a low frequency, after transformation with pMON9921 in which the *B.t.k.* coding sequence was expressed from the enhanced CaMV35S, promoter in pMON893, and in light of the fact that Elionor et al. (1989) report  
30 that the ats1A promoter itself is comparable in strength to the CaMV35S promoter and approximately 10-

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fold stronger when the CTP sequence is included. At least for the wild-type *B.t.k.* HD-1 coding sequence, this does not appear to be the case.

5 A variety of plant transformation vectors were constructed utilizing either the truncated synthetic HD-73 coding sequence of Figure 4 or the full length *B.t.k.* HD-73 coding sequence of Figure 11. These are listed in the table below.

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Table XV

5 Gene Constructs with CTPs

	<u>Vector</u>	<u>Promoter</u>	<u>CTP</u>	<u>B.t.k. HD-73</u>
				<u>Coding Sequence</u>
	pMON10806	En 35S	ats1A	truncated
	pMON10814	En35S	SSU11a	full length
10	pMON10811	SSU11a	SSU11a	truncated
	pMON10819	SSU11a	none	truncated
	pMON10815	ats1A	none	truncated
	pMON10817	ats1A	ats1A	truncated
	pMON10821	En 35S	ats1A	truncated
15	pMON10822	En 35S	ats1A	full length
	pMON10838	SSU11a	SSU11a	full length
	pMON10839	ats1A	ats1A	full length

All of the above vectors were used to transform tobacco plants. For all of the vectors containing truncated B.t.k. genes, leaf tissue from these plants has been analyzed for toxicity to insects and B.t.k. protein levels by immunoassay. pMON10806, 10811, 10819 and 10821 produce levels of B.t.k. protein comparable to pMON5383 and pMON5390 which contain synthetic B.t.k. HD-73 coding sequences driven by the En 35S promoter itself with no CTP. These plants also have the insecticidal activity expected for the B.t.k. protein levels detected. For pMON10815 and pMON10817 (containing the ats1A promoter), the level of B.t.k. protein is about 5-fold higher than that found in plants containing pMON5383 or 5390. These plants also

5 have higher insecticidal activity. Plants containing 10815 and 10817 contain up to 1% of their total soluble leaf protein as *B.t.k.* HD-73. This is the highest level of *B.t.k.* protein yet obtained with any of the synthetic genes.

10 This result is surprising in two respects. First, as noted above, the wild type coding sequences fused to the *ats1A* promoter and CTP did not show any evidence of higher levels of expression than for En 35S, and in fact had lower expression based on the absence of any insecticidal plants. Second, Elionor et al. (1989) show that for two other genes, the *ats1A* CTP can increase expression from the *ats1A* promoter by 15 about 10-fold. For the synthetic *B.t.k.* HD-73 gene, there is no consistent increase seen by including the CTP over and above that seen for the *ats1A* promoter alone.

20 Tobacco plants containing the full length synthetic HD-73 fused to the *SSU11A* CTP and driven by the En 35S promoter produced levels of *B.t.k.* protein and insecticidal activity comparable to pMON1518 which contains does not include the CTP. In addition, for 25 pMON10518 the *B.t.k.* protein extracted from plants was observed by gel electrophoresis to contain multiple forms less than full length, apparently due to the cleavage of the C-terminal portion (not required for toxicity) in the cytoplasm. For pMON10814, the majority of the protein appeared to be intact full length indicating that the protein has been stabilized 30 from proteolysis by targeting to the chloroplast.

Example 11 -- Targeting of B.t. Proteins to the Extracellular Space or Vacuole through the Use of Signal Peptides

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The B.t. proteins produced from the synthetic genes described here are localized to the cytoplasm of the plant cell, and this cytoplasmic localization results in plants that are insecticidally effective. It may 10 be advantageous for some purposes to direct the B.t. proteins to other compartments of the plant cell. Localizing B.t. proteins in compartments other than the cytoplasm may result in less exposure of the B.t. proteins to cytoplasmic proteases leading to greater 15 accumulation of the protein yielding enhanced insecticidal activity. Extracellular localization could lead to more efficient exposure of certain insects to the B.t. proteins leading to greater efficacy. If a B.t. protein were found to be 20 deleterious to plant cell function, then localization to a noncytoplasmic compartment could protect these cells from the protein.

In plants as well as other eucaryotes, proteins that are destined to be localized either extracellularly or in several specific compartments 25 are typically synthesized with an N-terminal amino acid extension known as the signal peptide. This signal peptide directs the protein to enter the compartmentalization pathway, and it is typically cleaved from the mature protein as an early step in 30 compartmentalization. For an extracellular protein, the secretory pathway typically involves

5 cotranslational insertion into the endoplasmic reticulum with cleavage of the signal peptide occurring at this stage. The mature protein then passes thru the Golgi body into vesicles that fuse with the plasma membrane thus releasing the protein into the extracellular space. Proteins destined for other compartments follow a similar pathway. For example, 10 proteins that are destined for the endoplasmic reticulum or the Golgi body follow this scheme, but they are specifically retained in the appropriate compartment. In plants, some proteins are also targeted to the vacuole, another membrane bound compartment in the cytoplasm of many plant cells. 15 Vacuole targeted proteins diverge from the above pathway at the Golgi body where they enter vesicles that fuse with the vacuole.

20 A common feature of this protein targeting is the signal peptide that initiates the compartmentalization process. Fusing a signal peptide to a protein will in many cases lead to the targeting of that protein to the endoplasmic reticulum. The efficiency of this step may depend on the sequence of the mature protein itself as well. The signals that direct a protein to 25 a specific compartment rather than to the extracellular space are not as clearly defined. It appears that many of the signals that direct the protein to specific compartments are contained within the amino acid sequence of the mature protein. This has been shown for some vacuole targeted proteins, but 30 it is not yet possible to define these sequences precisely. It appears that secretion into the

extracellular space is the "default" pathway for a protein that contains a signal sequence but no other compartmentalization signals. Thus, a strategy to direct *B.t.* proteins out of the cytoplasm is to fuse the genes for synthetic *B.t.* genes to DNA sequences encoding known plant signal peptides. These fusion genes will give rise to *B.t.* proteins that enter the secretory pathway, and lead to extracellular secretion or targeting to the vacuole or other compartments.

Signal sequences for several plant genes have been described. One such sequence is for the tobacco pathogenesis related protein PR1b described by Cornelissen et al. The PR1b protein is normally localized to the extracellular space. Another type of signal peptide is contained on seed storage proteins of legumes. These proteins are localized to the protein body of seeds, which is a vacuole like compartment found in seeds. A signal peptide DNA sequence for the beta subunit of the 7S storage protein of common bean (*Phaseolus vulgaris*), Pvub has been described by Doyle et al. Based on the published these published sequences, genes were synthesized by chemical synthesis of oligonucleotides that encoded the signal peptides for PR1b and Pvub. The synthetic genes for these signal peptides corresponded exactly to the reported DNA sequences. Just upstream of the translational initiation codon of each signal peptide a BamHI and BglII site were inserted with the BamHI site at the 5' end. This allowed the insertion of the signal peptide encoding segments into the BglII site

of pMON893 for expression from the En 35S promoter. In some cases to achieve secretion or compartmentalization of heterologous proteins, it has proved necessary to include some amino acid sequence beyond the normal cleavage site of the signal peptide. This may be necessary to insure proper cleavage of the signal peptide. For PR1b the synthetic DNA sequence also included the first 10 amino acids of mature PR1b. For PvuB the synthetic DNA sequence included the first 13 amino acids of mature PvuB. Both synthetic signal peptide encoding segments ended with NcoI sites to allow fusion in frame to the methionine initiation codon of the synthetic *B.t.* genes.

15 Four vectors encoding synthetic *B.t.k.* HD-73 genes were constructed containing these signal peptides. The synthetic truncated HD-73 gene from pMON5383 was fused with the signal peptide sequence of PvuB and incorporated into pMON893 to create pMON10827. The 20 synthetic truncated HD-73 gene from pMON5383 was also fused with the signal peptide sequence of PR1b to create pMON10824. The full length synthetic HD-73 gene from pMON10518 was fused with the signal peptide sequence of PvuB and incorporated into pMON893 to create pMON10828. The full length synthetic HD-73 25 gene from pMON10518 was also fused with the signal peptide sequence of PR1b and incorporated into pMON893 to create pMON10825.

These vectors were used to transform tobacco plants and the plants were assayed for expression of the 30 *B.t.k.* protein by Western blot analysis and for insecticidal efficacy. pMON10824 and pMON10827

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produced amounts of *B.t.k.* protein in leaf comparable to the truncated HD-73 vectors, pMON5383 and pMON5390. 5 pMON10825 and pMON10828 produced full length *B.t.k.* protein in amounts comparable to pMON10518. In all cases, the plants were insecticidally active against tobacco hornworm.

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## Claims:

5 1. In a method for improving the expression of a heterologous gene in plants by modifying the structural coding sequence of said gene, the improvement which comprises reducing the occurrence of polyadenylation signals selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA,  
10 ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

2. The method of Claim 1 further comprising the improvement of reducing the occurrence of ATTAA sequences within the structural coding sequence.

15 3. A method for modifying a wild-type structural gene sequence which encodes an insecticidal protein of *Bacillus thuringiensis* to enhance the expression of said protein in plants which comprises:

- 20 a) removing polyadenylation signals contained in said wild-type gene while retaining a sequence which encodes said protein; and
- b) removing ATTAA sequences contained in said wild-type gene while retaining a sequence which encodes said protein.

25 4. A method of Claim 3 further comprising the removal of self-complementary sequences and replacement of such sequences with nonself-complementary DNA comprising plant preferred codons while retaining a structural gene sequence encoding said protein.

30 5. A method of Claim 4 further comprising the use of plant preferred sequences in the removal of the polyadenylation signals and ATTAA sequences.

6. A method of Claim 3 in which the polyadenylation signals are selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, 5 ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

7. A method of Claim 4 in which the polyadenylation signals are selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, 10 ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

8. A method of Claim 5 in which the polyadenylation signals are selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, 15 ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

9. A method for modifying a wild-type structural gene sequence which encodes an insecticidal protein of *Bacillus thuringiensis* to enhance the expression of 20 said protein in plants which comprises:

- a) identifying regions within said sequence with greater than four consecutive adenine or thymine nucleotides;
- b) modifying the regions of step (a) which have two or more polyadenylation signals within a ten base sequence to remove said signals while maintaining a gene sequence which encodes said protein; and
- c) modifying the 15-30 base regions surrounding the regions of step (a) to remove major plant polyadenylation signals, consecutive sequences containing more than one minor polyadenylation 25 30

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signal and consecutive sequences containing more than one ATTTA sequence while maintaining a gene sequence which encodes said protein.

5 10. A method of Claim 9 in which the major plant polyadenylation signals are selected from the group consisting of AATAAA and AATAAT.

10 11. A method of Claim 10 in which the polyadenylation signals are selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

15 12. A method of Claim 11 further comprising the use of plant preferred sequences in the removal of polyadenylation signals and ATTTA sequences.

13. A structural gene which encodes an insecticidal protein of *Bacillus thuringiensis*, said gene being substantially devoid of polyadenylation signals and ATTTA sequences.

20 14. A structural gene of Claim 13 which is substantially devoid of polyadenylation signals selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

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15. A structural gene of Claim 13 which encodes an insecticidal protein of *B.t.k.* HD-1 having the sequence:

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30

1	ATGGCTATAGAAACTGGTTACACCCCAATCGATATTCCT	40
41	TGTCGCTAACGCAATTCTTTGAGTGAATTGTTCCCGG	80
81	TGCTGGATTGTGTTAGGACTAGTTGATATTATCTGGGGA	120
121	ATTTTGGTCCCTCTCAATGGGACGCATTCTTGTACAAA	160
161	TTGAACAGCTCATCAACCAGAGAATCGAAGAGATTCGCTAG	200
201	GAATCAAGCCATTCTAGATTAGAAGGACTAAGCAATCTT	240
241	TATCAAATTACGCAGAATCTTTAGAGAGTGGGAAGCAG	280
281	ATCCTACTAATCCAGCATTAAAGAGAAGAGATGCGTATTCA	320
321	ATTCAATGACATGAACAGTGCCCTTACAACCGCTATTCTT	360
361	CTTTTGCAAGTTCAAAATTATCAAGTTCTCTCCTCTCCG	400
401	TGTACGTTCAAGCTGCCAACCTCCACCTCTCAGTTTGAG	440
441	AGATGTTCAAGTGTGTTGGACAAAGGTGGGATTGATGCC	480
481	GCGACTATCAATAGTCGTTATAATGATTAACTAGGCTTA	520

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	521	TTGGCAACTATACAGATCATGCTGTACGCTGGTACAATAC	560
5	561	GGGATTAGAGCGTGTATGGGGACCGGATTCTAGAGATTGG	600
	601	ATCAGGTACAACCAGTTCAGAAGAGAGCTTACACTAAGT	640
	641	TATTAGATATCGTTCTCTATTCCGAACTATGATAGTAG	680
10			
	681	AACGTATCCAATTGAAACAGTTCCCAATTAACAAGAGAA	720
	721	ATTTATACAAACCCAGTATTAGAAAATTTGATGGTAGTT	760
15	761	TTCGAGGCTCGGCTCAGGGCATAGAAGGAAGTATTAGGAG	800
	801	TCCACATTTGATGGATATACTTAATAGTATAACCATCTAT	840
	841	ACGGATGCTCATAGAGGAGAATACTACTGGTCCGGTCACC	880
20			
	881	AGATCATGGCTTCTCCTGTAGGGTTTCGGGCCAGAATT	920
	921	CACTTTCCGCTATATGGAACTATGGGAAATGCAGCTCCA	960
25	961	CAACAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATA	1000
	1001	GAACATTATCGTCCACCTTATATAGAAGACCTTTAACAT	1040
	1041	CGGGATCAACAACCAACAACATCTGTTCTTGACGGGACA	1080
30	1081	GAATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTG	1120

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1121	TATACAGAAAAAGCGGAACGGTAGATTGCTGGATGAAAT	1160
5		
1161	ACCGCCACAGAATAACAACGTGCCACCTAGGCAAGGATT	1200
1201	AGTCATCGATTAAGCCATGTTCAATGTTCGTTAGGCT	1240
1241	TTAGTAATAGTAGTGTAAGTATAATAAGAGCTCCTATGTT	1280
10		
1281	CTCTGGATAACATCGTAGTGCTGAGTTCAACAAACATCATC	1320
1321	CCTTCATCACAAATCACCCAAATCCCACTCACCAAGTCTA	1360
15	1361 CTAATCTGGCTCTGGAACCTCTGTCGTTAAAGGACCAGG	1400
1401	ATTTACAGGAGGAGATATTCTCGAAGAACCTCACCTGGC	1440
1441	CAGATTTAACCTTAAGAGTAAATATTACTGCACCATTAT	1480
20		
1481	CACAAAGATATCGGGTAAGAATTGCTACGCTTCTACCAC	1520
1521	AAACCTTCAGTCCACACATCAATTGACGGAAGACCTATT	1560
25	1561 AATCAGGGGAATTTTCAGCAACTATGAGTAGTGGGAGTA	1600
1601	ATTTACAGTCCGGAAGCTTCTAGGACTGTAGGTTTACTAC	1640
1641	TCCGTTAACCTTCAAATGGATCAAGTGTATTACGTTA	1680

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1681 AGTGCTCATGTCTCAATTCAAGGCAATGAAGTTATATAG 1720  
5 1721 ATCGAATTGAATTGTTCCGGCA 1743.

16. A structural gene of Claim 13 which encodes an insecticidal protein of B.t.k. HD-73 having the sequence:

10 1 ATGGCCATTGAAACCGGTTACACTCCCATCGACATCTCCT 40  
41 TGTCCCTTGACACAGTTCTGCTCAGCGAGTTCCGTGCCAGG 80  
15 81 TGCTGGGTTCGTTCTCGGACTAGTTGACATCATCTGGGGT 120  
121 ATCTTGGTCCATCTCAATGGGATGCATTCTGGTGCAAA 160  
161 TTGAGCAGTTGATCAACCAGAGGATCGAAGAGAGTTGCCAG 200  
20 201 GAACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTC 240  
241 TACCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCG 280  
25 281 ATCCTACTAACCCAGCTCTCCCGAGGAAATGCGTATTCA 320  
321 ATTCAACGACATGAACAGCGCCTTGACCACAGCTATCCCA 360  
361 TTGTTCGCAGTCCAGAACTACCAAGTTCTCTCTGTCCG 400  
30 401 TGTACGTTCAAGCAGCTAATCTCACCTCAGCGTGCTTCG 440

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	441	AGACGTTAGCGTGTGTTGGCAAAGGTGGGATTCGATGCT	480
5	481	GCAACCATCAATAGCCGTTACAACGACCTTACTAGGCTGA	520
	521	TTGGAAACTACACCGACCACGCTGTTGTTGACAACAC	560
	561	TGGCTTGGAGCGTGTCTGGGGCCTGATTCTAGAGATTGG	600
10			
	601	ATTAGATACAACCAGTTCAGGAGAGAATTGACCCTCACAG	640
	641	TTTTGGACATTGTGTCTCTTCCGAACATATGACTCCAG	680
15	681	AACCTACCCTATCCGTACAGTGTCCCAACTTACCAAGAGAA	720
	721	ATCTATACTAACCCAGTTCTTGAGAACTTCGACGGTAGCT	760
	761	TCCGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAG	800
20			
	801	CCCACACTTGATGGACATCTGAACAGCATAACTATCTAC	840
	841	ACCGATGCTCACAGAGGAGAGTATTACTGGTCTGGACACC	880
	881	AGATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTT	920
25			
	921	TACCTTCCTCTATGAACTATGGAAACGCCGCTCCA	960
	961	CAACAAACGTATCGTTGCTCAACTAGGTCAGGGTGTCTACA	1000
30	1001	GAACCTTGTCTTCCACCTTGTACAGAAGACCCTCAATAT	1040

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1041	CGGTATCAACAACCAGCAACTTCCGTTCTTGACGGAACA	1080
5	5 1081 GAGTCGCCTATGGAACCTCTTCTAACCTGCCATCCGCTG	1120
	1121 TTTACAGAAAGAGCGGAACCGTTGATTCTTGGACGAAAT	1160
	1161 CCCACCACAGAACACAATGTGCCACCCAGGCAAGGATTC	1200
10	10 1201 TCCCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGAT	1240
	1241 TCAGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTT	1280
15	15 1281 CTCTTGGATAACCGTAGTGCTGAGTTCAACAAACATCATC	1320
	1321 GCATCCGATAGTATTACTCAAATCCCTGCAGTGAAGGGAA	1360
	1361 ACTTTCTCTTCAACGGTTCTGTCATTCAAGGACCAGGATT	1400
20	20 1401 CACTGGTGGAGACCTCGTTAGACTCAACAGCAGTGGAAAT	1440
	1441 AACATTCAAGAATAGAGGGTATATTGAAGTTCCAATTCACT	1480
	1481 TCCCACATCACATCTACCAAGATATAGAGTTCGTGTGAGGTA	1520
25	25 1521 TGCTTCTGTGACCCCTATTCAACCTCAACGTTAATTGGGGT	1560
	1561 AATTCAATCCATCTCTCCAATACAGTTCCAGCTACAGCTA	1600
30	30 1601 CCTCCTTGGATAATCTCCAATCCAGCGATTCGGTTACTT	1640

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1641 TGAAAGTGCCAATGCTTTACATCTTCACTCGGTAAACATC 1680  
5 1681 GTGGGTGTTAGAAACTTAGTGGGACTGCAGGAGTGATTA 1720  
1721 TCGACAGATTGAGTTCATTCCAGTTACTGCAACACTCGA 1760  
10 1761 GGCTGAG 1767.

17. A structural gene of Claim 13 encoding a insecticidal protein of *B.t.k.* HD-1 having the sequence:

15 1 ATGGACAACAAACCCAAACATCAACGAATGCATTCCATACA 40  
41 ACTGCTTGAGTAACCCAGAAGTTGAAGTACTTGGTGGAGA 80  
81 ACGCATTGAAACCGGTTACACTCCCATCGACATCTCCTTG 120  
20 121 TCCTTGACACAGTTCTGCTCAGCGAGTTCGTGCCAGGTG 160  
161 CTGGGTTCGTCTCGGACTAGTTGACATCATCTGGGTAT 200  
25 201 CTTTGGTCCATCTCAATGGGATGCATTCTGGTGCAAATT 240  
241 GAGCAGTTGATCAACCAGAGGGATCGAAGAGTTGCCAGGA 280  
281 ACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTCTA 320  
30 321 CCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCGAT 360

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361	CCTACTAACCCAGCTCTCCGCGAGGAAATGCGTATTCAAT	400
5	401 TCAACGACATGAACAGCGCCTTGACCACAGCTATCCCATT	440
441	GTTCGCAGTCCAGAACTACCAAGTT CCTCTTGTCCGTG	480
10	481 TACGTTCAAGCAGCTAATCTCACCTCAGCGTGCTTCGAG	520
521	ACGTTAGCGTGTGGGCAAAGGTGGGGATTCGATGCTGC	560
561	AACCATCAATAGCCGTTACAACGACCTTACTAGGCTGATT	600
15	601 GGAAACTACACCGACCACGCTGTTGTTGGTACAACACTG	640
641	GCTTGGAGCGTGTCTGGGTCTGATTCTAGAGATTGGAT	680
681	TAGATACAACCAGTTCAGGAGAGAATTGACCCACAGTT	720
20	721 TTGGACATTGTGTCTCTTCCCAGACTATGACTCCAGAA	760
761	CCTACCCCTATCCGTACAGTGTCCCAACTTACCAAGAGAAAT	800
25	801 CTATACTAACCCAGTTCTTGAGAACTTCGACGGTAGCTC	840
841	CGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAGCC	880
881	CACACTTGATGGACATCTTGAACAGCATAACTATCTACAC	920
30	921 CGATGCTCACAGAGGAGAGTATTACTGGTCTGGACACCAG	960

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961	ATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTTA	1000
5	1001 CCTTTCCCTCTCTATGGAACATATGGGAAACGCCGCTCCACA	1040
	1041 ACAACGTATCGTTGCTCAACTAGGTCAAGGGTGTCTACAGA	1080
10	1081 ACCTTGTCTTCCACCTTGTACAGAAGACCCTTCAATATCG	1120
	1121 GTATCAACAACCAGCAACTTCCGTTCTTACAGGAAACAGA	1160
	1161 GTTCGCCTATGGAACCTCTTCTAACTGCCATCCGCTGTT	1200
15	1201 TACAGAAAGAGCGGAACCGTTGATTCCCTGGACGAAATCC	1240
	1241 CACCACAGAACAAACAATGTGCCACCCAGGCAAGGATTCTC	1280
20	1281 CCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGATTTC	1320
	1321 AGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTTCT	1360
	1361 CATGGATTCATCGTAGTGCTGAGTTCAACAATATCATTCC	1400
25	1401 TTCCTCTCAAATCACCCAAATCCCATTGACCAAGTCTACT	1440
	1441 AACCTTGGATCTGGAACCTCTGTGAAAGGACCAAGGCT	1480
	1481 TCACAGGAGGTGATATTCTTAGAAGAACCTCTCCTGGCCA	1520
30	1521 GATTAGCACCCCTCAGAGTTAACATCACTGCACCACCTTCT	1560

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1561	CAAAGATATCGTGTCAAGGATTGTTACGCATCTACCACTA	1600	
5	1601	ACTTGCAATTCCACACCTCCATCGACGGAAGGCCTATCAA	1640
	1641	TCAGGGTAACCTCTCCGCAACCATGTCAAGCGGCAGCAAC	1680
	1681	TTGCAATCCGGCAGCTTCAGAACCGTCGGTTCACTACTC	1720
10	1721	CTTCAACTCTCTAACGGATCAAGCGTTTCACCCCTAG	1760
	1761	CGCTCATGTGTTCAATTCTGGCAATGAAGTGTACATTGAC	1800
	1801	CGTATTGAGTTGTGCCCTGCCGAAGTTACCTCGAGGCTG	1840
	1841	AGTAC 1845.	

18. A structural gene of Claim 13 encoding an  
20 insecticidal protein derived from B.t.k. HD-73 having  
the sequence:

1	ATGGACACAAACCCAAACATCAACGAATGCATTCCATACA	40	
25	41	ACTGCTTGAGTAACCCAGAACGTTGAAGTACTTGGTGGAGA	80
	81	ACGCATTGAAACCGGTTACACTCCCATCGACATCTCCTTG	120
	121	TCCTTGACACAGTTCTGCTCAGCGAGTTCGTGCCAGGTG	160
30	161	CTGGGTTCGTTCTCGGACTAGTTGACATCATCTGGGTAT	200

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201	CTTTGGTCCATCTCAATGGGATGCATTCTGGTGCAAATT	240	
5	5	GAGCAGTTGATCAACCAGAGGATCGAAGAGAGTTGCCAGGA	280
281	ACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTCTA	320	
321	321 CCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCGAT	360	
10	361 CCTACTAACCCAGCTCTCCGCGAGGAAATGCGTATTCAAT	400	
401	401 TCAACGACATGAACAGCGCCTTGACCACAGCTATCCCATT	440	
15	441 GTTCGCAGTCCAGAACTACCAAGTTCTCTCTTGCCGTG	480	
481	481 TACGTTCAAGCAGCTAATCTCACCTCAGCGTGTTCGAG	520	
521	521 ACGTTAGCGTGTGGCAAAAGGTGGGATTGATGCTGC	560	
20	561 AACCATCAATAGCCGTTACAACGACCTTACTAGGCTGATT	600	
601	601 GGAAACTACACCGACCACGCTGTTGTTGGTACAACACTG	640	
641	641 GCTTGGAGCGTGTCTGGGTCTGATTCTAGAGATTGGAT	680	
25	681 TAGATACAACCAGTCAGGAGAGAATTGACCCACAGTT	720	
721	721 TTGGACATTGTGTCTCTTCCGAACATGACTCCAGAA	760	
30	761 CCTACCCTATCCGTACAGTGTCCCAACTTACCAAGAGAAAT	800	

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801	CTATACTAACCCAGTTCTTGAGAACCTCGACGGTAGCTTC	840
5	CGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAGCC	880
881	CACACTTGATGGACATCTTGAACAGCATAACTATCTACAC	920
921	CGATGCTCACAGAGGGAGAGTATTACTGGTCTGGACACCAG	960
10	ATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTTA	1000
1001	CCTTCCTCTATGGAACTATGGAAACGCCCCTCCACA	1040
15	ACAACGTATCGTTGCTCAACTAGGTCAAGGTGTCTACAGA	1080
1081	ACCTTGTCTTCCACCTTGTACAGAAGACCCCTCAATATCG	1120
1121	GTATCAACAAACCAACAGCAACTTCCGTTCTGACGGAACAGA	1160
20	GTTGCCTATGGAACCTCTTCTAACTTGCATCCGCTGTT	1200
1201	TACAGAAAGAGCGGAACCGTTGATTCTTGGACGAAATCC	1240
25	CACCACAGAACAAACAATGTGCCACCCAGGCAAGGATTCTC	1280
1281	CCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGATTTC	1320
1321	AGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTTCT	1360
30	CTTGGATACACCGTAGTGCTGAGTTCAACAAACATCATCGC	1400

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1401	ATCCGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC	1440
5	1441 TTTCTCTCAACGGTTCTGTCATTCAGGACCAGGATTCA	1480
1481	CTGGTGGAGACCTCGTTAGACTCAACAGCAGTGGAAATAA	1520
10	1521 CATTAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
1561	CCATCCACATCTACCAAGATATAGAGTTCGTGTGAGGTATG	1600
1601	CTTCTGTGACCCCTATTCAACCTCAACGTTAATTGGGTAA	1640
15	1641 TTCATCCATCTTCTCCAATACAGTTCCAGCTACAGCTACC	1680
1681	TCCTTGGATAATCTCCAATCCAGCGATTCGGTTACTTTG	1720
1721	AAAGTGCCAATGCTTTACATCTTCACTCGGTAACATCGT	1760
20	1761 GGGTGTAGAAACTTTAGTGGGACTGCAGGAGTGATTATC	1800
1801	GACAGATTCGAGTTCAATTCCAGTTACTGCAACACTCGAGG	1840
25	1841 CTGAATATAATCTGGAAAGAGCGCAGAAGGCGGTAAATGCG	1880
1881	CTGTTACGTCTACAAACCAGCTTGGACTCAAGACAAATG	1920.

19. A structural gene of Claim 13 encoding the  
full-length insecticidal protein of *B.t.k.* HD-73  
30 having the sequence:

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1	ATGGACAACAACCAAACATCAACGAATGCATTCCATACA	40
5	ACTGCTTGAGTAACCCAGAAGTTGAAGTACTTGGTGGAGA	80
81	ACGCATTGAAACCGGTTACACTCCCATCGACATCTCCTTG	120
121	TCCTTGACACAGTTCTGCTCAGCGAGTTCGTGCCAGGTG	160
10	CTGGGTTCGTTCTCGGACTAGTTGACATCATCTGGGTAT	200
161	CTTTGGTCCATCTCAATGGGATGCATTCTGGTGCAAATT	240
15	241 GAGCAGTTGATCAACCAGAGGATCGAAGAGTTCGCCAGGA	280
281	ACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTCTA	320
321	CCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCGAT	360
20	361 CCTACTAACCCAGCTCTCCCGAGGAAATGCGTATTCAAT	400
401	TCAACGACATGAACAGCGCCTTGACCACAGCTATCCCATT	440
25	441 GTTCGCAGTCCAGAACTACCAAGTTCCCTCTCTGTCCGTG	480
481	TACGTTCAAGCAGCTAATCTCACCTCAGCGTGCTTCGAG	520
521	ACGTTAGCGTGTGAAAGGTGGGATTGATGCTGC	560
30	561 AACCATCAATAGCCGTTACAACGACCTTACTAGGCTGATT	600

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	601	GGAAACTACACCGACCACGCTGTTGTTGGTACAACACTG	640
5	641	GCTTGGAGCGTGTCTGGGGCCTGATTCTAGAGATTGGAT	680
	681	TAGATACAACCAGTTCAGGAGAGAATTGACCCTCACAGTT	720
	721	TTGGACATTGTGTCTCTCTTCCGAACACTATGACTCCAGAA	760
10	761	CCTACCCCTATCCGTACAGTGTCCCAACTTACCGAGAGAAAT	800
	801	CTATACTAACCCAGTTCTTGAGAACCTTCGACGGTAGCTTC	840
15	841	CGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAGCC	880
	881	CACACTTGATGGACATCTTGAACAGCATAACTATCTACAC	920
	921	CGATGCTCACAGAGGAGAGTATTACTGGTCTGGACACCAG	960
20	961	ATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTTA	1000
	1001	CCTTCCTCTATGGAACATATGGGAAACGCCGCTCCACA	1040
	1041	ACAACGTATCGTTGCTCAACTAGGTCAAGGGTGTCTACAGA	1080
25	1081	ACCTTGTCTTCCACCTTGTACAGAACCCCTCAATATCG	1120
	1121	GTATCAACAACCAGCAACTTCCGTTCTGACGGAACAGA	1160
30	1161	GTTCGCCTATGGAACCTCTTAACCTGCCATCCGCTGTT	1200

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1201	TACAGAAAGAGCGGAACCGTTGATTCCCTGGACGAAATCC	1240
5	• • •	• • •
1241	CACCAAGAACAAACAATGTGCCACCCAGGCAAGGATTCTC	1280
•	• • •	• • •
1281	CCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGATTC	1320
•	• • •	• • •
1321	AGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTTCT	1360
10	• • •	• • •
1361	CTTGGATACACCGTAGTGCTGAGTTCAACAAACATCATCGC	1400
•	• • •	• • •
1401	ATCCGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC	1440
•	• • •	• • •
15	1441 TTTCTCTTCAACGGTTCTGTCATTCAGGACCAGGATTCA	1480
•	• • •	• • •
1481	CTGGTGGAGACCTCGTTAGACTCAACAGCAGTGGAAATAA	1520
•	• • •	• • •
1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
20	• • •	• • •
1561	CCATCCACATCTACCAGATATAGAGTTCGTGTGAGGTATG	1600
•	• • •	• • •
1601	CTTCTGTGACCCCTATTCACCTCAACGTTAATTGGGGTAA	1640
•	• • •	• • •
1641	TTCATCCATCTTCTCCAATACAGTTCCAGCTACAGCTACC	1680
25	• • •	• • •
1681	TCCTTGGATAATCTCCAATCCAGCGATTCGGTTACTTTG	1720
•	• • •	• • •
1721	AAAGTGCCAATGCTTTACATCTTCACTCGGTAACATCGT	1760
•	• • •	• • •
30	1761 GGGTGTAGAAACTTTAGTGGGACTGCAGGAGTGATTATC	1800

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	1801	GACAGATTGAGTTCATTCAGTTACTGCAACACTCGAGG	1840
5	1841	CTGAATATAATCTGGAAAGAGCGCAGAAGGCCGTGAATGC	1880
	1881	GCTGTTACGTCTACAAACCAGCTCGGCCTCAAGACCAAT	1920
	1921	GTGACGGATTATCATATTGATCAAGTGTCCAACTTGGTGA	1960
10	1961	CCTACCTCAGCGATGAGTTCTGTCTGGATGAAAAGCGAGA	2000
	2001	ATTGTCCGAGAAAGTCAAACATGCGAAGCGACTCAGTGAT	2040
15	2041	GAACGCAATTTACTCCAAGATTCAAATTCAAAGACATTA	2080
	2081	ATAGGCAACCAGAACGTGGGTGGGGCGGAAGTACAGGGAT	2120
	2121	TACCATCCAGGGAGGTGACGACGTGTTCAAGGAGAACTAC	2160
20	2161	GTCACACTATCAGGTACCTTGATGAGTGCTATCCAACAT	2200
	2201	ACCTCTACCAGAAGATCGACGAGTCCAAGTTGAAAGCCTT	2240
25	2241	TACCCGTTATCAATTAAAGAGGGTATATCGAAGATAGTCAA	2280
	2281	GACCTCGAGATCTACCTCATCCGCTACAATGCAAAACATG	2320
	2321	AAACAGTAAATGTGCCAGGTACGGGTTCTTATGGCCGCT	2360
30	2361	TTCAGCCAAAGTCCAATCGGAAAGTGTGGAGAGCCGAAT	2400

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	2401	CGATGCGCGCCACACCTGAATGGAATCCTGACTTAGATT	2440
5	2441	GTTCGTGTAGGGATGGAGAAAAGTGTGCCCATTCGCA	2480
	2481	TCATTTCTCCTTAGACATTGATGTAGGATGTACAGACTTA	2520
	2521	AATGAGGGACCTAGGTGTATGGGTGATCTTAAGATTAAGA	2560
10			
	2561	CGCAAGATGGGCACGCAAGACTAGGGATCTAGAGTTCT	2600
	2601	CGAAGAGAAACCATTAGTAGGAGAAGCGCTAGCTCGTG	2640
15	2641	AAAAGAGCGGAGAAAAATGGAGAGACAAACGTGAGAAGT	2680
	2681	TGGAATGGGAGACCAACATCGTCTACAAAGAGGCAAAAGA	2720
	2721	ATCTGTAGATGCTTATTGTAACTCTCAATATGATCAA	2760
20			
	2761	TTACAAGCGGATACGAATATTGCCATGATTGCGGCAG	2800
	2801	ATAAACGTGTTCATAGCATTGAGAAGCTTATCTGCCTGA	2840
25	2841	GCTGTCTGTGATTCCGGGTGTCAATGCGGCTATTTTGAA	2880
	2881	GAATTAGAAGGGCGTATTTCACTGCATTCTCCCTCTACG	2920
	2921	ATGCCAGAAACGTCATCAAGAACGGTGACTTCAACAATGG	2960
30	2961	CTTATCCTGCTGAAACGTGAAAGGGCATGTAGATGTAGAA	3000

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	3001	GAACAAAACAACCAACGTTGGTCCTTGTGTTCCGGAAT	3040
5	3041	GGGAAGCAGAAGTGTACAAGAAGTTCGTGTCTGTCCGGG	3080
	3081	TCGTGGCTATATCCTCGTGTACAGCGTACAAGGAGGG	3120
10	3121	TATGGAGAAGGTTGCGTAACCATTGAGATCGAGAACAA	3160
	3161	ATACAGACGAACTGAAGTTAGCAACTGCGTAGAAGAGGA	3200
	3201	AATCTATCCAAAATAACACCGTAACGTGTAATGATTAACT	3240
15	3241	GTAAATCAAGAAGAATACGGAGGTGCGTACACTCTCGTA	3280
	3281	ATCGAGGATATAACGAAGCTCCTCCGTACCGAGCTGATTA	3320
20	3321	TGCGTCAGTCTATGAAGAAAATCGTATACAGATGGACGA	3360
	3361	AGAGAGAATCCTTGTGAATTAAACAGAGGGTATAGGGATT	3400
	3401	ACACGCCACTACCAGTTGGTTATGTGACAAAAGAATTAGA	3440
25	3441	ATACTTCCCAGAAACCGATAAGGTATGGATTGAGATTGGA	3480
	3481	GAAACGGAAGGAACATTATCGTGGACAGCGTGGATTAC	3520
	3521	TCCTTATGGAGGAA 3534.	

30

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20. A structural gene of Claim 13 encoding a full-length insecticidal protein of *B.t.k.* HD-73 having the sequence:

5

1	ATGGACAACAACCCAAACATCAACGAATGCATTCCATACA	40
41	ACTGCTTGAGTAACCCAGAAGTTGAAGTACTTGGTGGAGA	80
10	81 ACGCATTGAAACCGGTTACACTCCCATCGACATCTCCTTG	120
	121 TCCTTGACACAGTTCTGCTCAGCGAGTTCGTGCCAGGTG	160
	161 CTGGGTTCGTTCTCGGACTAGTTGACATCATCTGGGTAT	200
15	201 CTTTGGTCCATCTCAATGGGATGCATTCTGGTGCAAATT	240
	241 GAGCAGTTGATCAACCAGAGGATCGAAGAGAGTTGCCAGGA	280
20	281 ACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTCTA	320
	321 CCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCGAT	360
	361 CCTACTAACCCAGCTCTCGCGAGGAAATGCGTATTCAAT	400
25	401 TCAACGACATGAACAGCGCCTTGACCACAGCTATCCCATT	440
	441 GTTCCGCAGTCCAGAACTACCAAGTTCTCTCTTGCCGTG	480
30	481 TACGTTCAAGCAGCTAATCTCACCTCAGCGTGCTCGAG	520

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	521	ACGTTAGCGTGGGCAAAGGTGGGATTGATGCTGC	560
5	561	AACCATAAGCCGTTACAACGACCTTACTAGGCTGATT	600
	601	GGAAACTACACCGACCACGCTGTTGTTGACAACACTG	640
10	641	GCTTGGAGCGTGTCTGGGGCTGATTCTAGAGATTGGAT	680
	681	TAGATACAACCAGTTCAGGAGAGAATTGACCCCTCACAGTT	720
	721	TTGGACATTGTGTCTCTTCCCAGACTATGACTCCAGAA	760
15	761	CCTACCCTATCCGTACAGTGTCCAACTTACCAAGAGAAAT	800
	801	CTATACTAACCCAGTTCTTGAGAACCTCGACGGTAGCTTC	840
	841	CGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAGCC	880
20	881	CACACTTGATGGACATCTTGAACAGCATAACTATCTACAC	920
	921	CGATGCTCACAGAGGAGAGTATTACTGGTCTGGACACCAG	960
25	961	ATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTTA	1000
	1001	CCTTCCTCTATGGAACATATGGAAACGCCGCTCCACA	1040
	1041	ACAACGTATCGTTGCTCAACTAGGTCAAGGGTCTACAGA	1080
30	1081	ACCTTGTCTTCCACCTTGTACAGAAGACCCTCAATATCG	1120

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	1121	GTATCAACAACCAGCAACTTCCGTTCTGACGGAACAGA	1160
5	1161	GTTCGCCTATGGAACCTCTTCTAACCTGCCATCCGCTGTT	1200
	1201	TACAGAAAGAGCGGAACCGTTGATTCCCTGGACGAAATCC	1240
	1241	CACCAACAGAACAAACATGTGCCACCCAGGCAAGGATTCTC	1280
10		.	.
	1281	CCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGATTTC	1320
	1321	AGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTTCT	1360
15	1361	CTTGGATACACCGTAGTGCTGAGTTCAACAAACATCATCGC	1400
	1401	ATCCGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC	1440
	1441	TTTCTCTTCAACGGTTCTGTCATTCAGGACCAGGATTCA	1480
20		.	.
	1481	CTGGTGGAGACCTCGTTAGACTCAACAGCAGTGGAAATAA	1520
	1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
	1561	CCATCCACATCTACCAGATATAGAGTTCGTGTGAGGTATG	1600
25		.	.
	1601	CTTCTGTGACCCCTATTCACCTCAACGTTAATTGGGGTAA	1640
	1641	TTCATCCATCTTCTCCAATACAGTTCCAGCTACAGCTACC	1680
	1681	TCCTTGGATAATCTCCAATCCAGCGATTCGGTTACTTTG	1720

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	1721	AAAGTGCCAATGCTTTACATCTTCACTCGGTAACATCGT	1760
5	1761	GGGTGTTAGAAACTTTAGTGGGACTGCAGGAGTGATTATC	1800
	1801	GACAGATTGAGTTCATTCCAGTTACTGCAACACTCGAGG	1840
	1841	CTGAATATAATCTGGAAAGAGCGCAGAAGGCCGTGAATGC	1880
10		.	.
	1881	GCTGTTACGTCTACAAACCAACTAGGGCTAAAAACAAAT	1920
	1921	GTAACGGATTATCATATTGATCAAGTGTCCAATTTAGTTA	1960
15	1961	CGTATTATCGGATGAATTTGTCTGGATGAAAAGCGAGA	2000
	2001	ATTGTCCGAGAAAGTCAAACATGCGAAGCGACTCAGTGAT	2040
	2041	GAACGCAATTACTCCAAGATTCAAATTCAAAGACATTA	2080
20		.	.
	2081	ATAGGCAACCAGAACGTGGGTGGGGCGGAAGTACAGGGAT	2120
	2121	TACCATCCAAGGAGGGATGACGTATTTAAAGAAAATTAC	2160
25	2161	GTCACACTATCAGGTACCTTGATGAGTGCTATCCAACAT	2200
	2201	ATTTGTATCAAAAAATCGATGAATCAAATTAAAAGCCTT	2240
	2241	TACCCGTTATCAATTAAGAGGGTATATCGAAGATAGTCAA	2280
30	2281	GACTTAGAAATCTATTAATTGCTACAATGCAAAACATG	2320

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	2321	AAACAGTAAATGTGCCAGGTACGGGTTCCCTATGGCCGCT	2360
5	2361	TTCAGCCCAAAGTCCAATCGAAAGTGTGGAGAGCCGAAT	2400
	2401	CGATGCGCGCCACACCTTGAATGGAATCCTGACTTAGATT	2440
	2441	GTTCGTGTAGGGATGGAGAAAAGTGTGCCCATTCGCA	2480
10			
	2481	TCATTTCTCCTTAGACATTGATGTAGGATGTACAGACTTA	2520
	2521	AATGAGGACCTAGGTGTATGGGTGATCTTAAGATTAAGA	2560
15	2561	CGCAAGATGGCACGCAAGACTAGGAAATCTAGAGTTCT	2600
	2601	CGAAGAGAAACCATTAGTAGGAGAAGCGCTAGCTCGTGTG	2640
	2641	AAAAGAGCGGAGAAAAATGGAGAGACAAACGTGAAAAT	2680
20			
	2681	TGGAATGGAAACAAATATCGTTATAAAGAGGGCAAAGA	2720
	2721	ATCTGTAGATGCTTATTGTAAACTCTCAATATGATCAA	2760
25	2761	TTACAAGCGGATACGAATATTGCCATGATTGCGGCAG	2800
	2801	ATAAACGTGTTCATAGCATTGAGAAGCTTATCTGCCTGA	2840
	2841	GCTGTCTGTGATTCCGGGTGTCAATGCGGCTATTTTGAA	2880
30	2881	GAATTAGAAGGGCGTATTTCACTGCATTCTCCCTATATG	2920

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	2921	ATGCGAGAAATGTCATTAAAATGGTGATTTAATAATGG	2960
5	2961	CTTATCCTGCTGGAACGTGAAAGGGCATGTAGATGTAGAA	3000
	3001	GAACAAAACAACCAACGTTGGTCCTTGTGTTCCCGGAAT	3040
	3041	GGGAAGCAGAAGTGTACAAGAAGTTCGTGTCTGTCCGGG	3080
10		.	.
	3081	TCGTGGCTATATCCTTCGTGTACAGCGTACAAGGAGGGA	3120
	3121	TATGGAGAAGGTTGCGTAACCATTGAGATCGAGAACAA	3160
15	3161	ATACAGACGAACGTAAAGTTAGCAAACGCGTAGAAGAGGA	3200
	3201	AATCTATCCAAATAACACGGTAACGTGTAATGATTATACT	3240
	3241	GTAAATCAAGAAGAATAACGGAGGTGCGTACACTCTCGTA	3280
20		.	.
	3281	ATCGAGGATATAACGAAGCTCCTCCGTACCGAGCTGATTA	3320
	3321	TGCGTCAGTCTATGAAGAAAATCGTATAACAGATGGACGA	3360
	3361	AGAGAGAATCCTTGTGAATTAAACAGAGGGTATAGGGATT	3400
25		.	.
	3401	ACACGCCACTACCAGTTGGTTATGTGACAAAAGAATTAGA	3440
	3441	ATACTTCCCAGAAACCGATAAGGTATGGATTGAGATTGGA	3480
30			

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3481 GAAACGGAAGGAACATTTATCGTGGACAGCGTGGATTAC 3520  
5 3521 TCCTTATGGAGGAA 3534.

21. A structural gene of Claim 13 encoding a full-length insecticidal protein of *B.t.k.* HD-73 having the sequence:

10 1 ATGGACAACAACCCAAACATCAACGAATGCATTCCATACA 40  
41 ACTGCTTGAGTAACCCAGAAGTTGAAGTACTTGGTGGAGA 80  
15 81 ACGCATTGAAACCGGTTACACTCCCATCGACATCTCCTTG 120  
121 TCCTTGACACAGTTCTGCTCAGCGAGTTCGTGCCAGGTG 160  
161 CTGGGTTCGTTCTCGGACTAGTTGACATCATCTGGGTAT 200  
20 201 CTTTGGTCCATCTCAATGGGATGCATTCCCTGGTGCAAATT 240  
241 GAGCAGTTGATCAACCAGAGGATCGAAGAGTTCCGCCAGGA 280  
25 281 ACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTCTA 320  
321 CCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCGAT 360  
361 CCTACTAACCCAGCTCTCCGCGAGGAAATGCGTATTCAAT 400  
30 401 TCAACGACATGAACAGCGCCTTGACCACAGCTATCCCATT 440

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441	GTTCGCAGTCCAGAACTACCAAGTTCCCTCTTGTCCGTG	480	
5	5	481 TACGTTCAAGCAGCTAATCTCACCTCAGCGTGCTCGAG	520
521	ACGTTAGCGTGTGTTGGGCAAAGGTGGGGATTCGATGCTGC	560	
10	561 AACCATCAATAGCCGTTACAACGACCTTACTAGGCTGATT	600	
601	GGAAACTACACCGACCACGCTGTTGTTGGTACAACACTG	640	
641	GCTTGGAGCGTGTCTGGGGCCTGATTCTAGAGATTGGAT	680	
15	681 TAGATACAACCAGTTCAAGGAGAGAATTGACCCTCACAGTT	720	
721	TTGGACATTGTGTCTCTTCCCGAACTATGACTCCAGAA	760	
20	761 CCTACCCTATCCGTACAGTGTCCCAACTTACCAAGAGAAAT	800	
801	CTATACTAACCCAGTTCTTGAGAACTTCGACGGTAGCTTC	840	
841	CGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAGCC	880	
25	881 CACACTTGATGGACATCTGAACAGCATAACTATCTACAC	920	
921	CGATGCTCACAGAGGGAGAGTATTACTGGTCTGGACACCAG	960	
961	ATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTTA	1000	
30	1001 CCTTTCTCTATGAACTATGGAAACGCCGCTCCACA	1040	

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1041	ACAAACGTATCGTTGCTCAACTAGGTCAAGGGTGTCTACAGA	1080
5	ACCTTGCTTCCACCTTGTACAGAAGACCCCTCAATATCG	1120
1121	GTATCAACAAACCAGCAACTTCCGTTCTGACGGAACAGA	1160
10	GTTCGCCATGGAACCTCTCTAACTTGCCATCCGCTGTT	1200
1201	TACAGAAAGAGCGGAACCGTTGATTCTTGGACGAAATCC	1240
1241	CACCACAGAACAAACAATGTGCCACCCAGGCAAGGATTCTC	1280
15	CCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGATT	1320
1321	AGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTTCT	1360
1361	CTTGGATACACCGTAGTGCTGAGTTCAACAAACATCATCGC	1400
20	ATCCGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC	1440
1441	TTTCTCTCAACGGTTCTGTCAATTTCAGGACCAGGATTCA	1480
25	CTGGTGGAGACCTCGTTAGACTCAACAGCAGTGGAAATAA	1520
1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
1561	CCATCCACATCTACCAGATATAGAGTTCGTGTGAGGTATG	1600
30	CTTCTGTGACCCCTATTCACCTAACGTTAATTGGGGTAA	1640

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	1641	TTCATCCATCTCTCCAATACAGTTCCAGCTACAGCTACC	1680
5	1681	TCCTTGGATAATCTCCAATCCAGCGATTCGGTTACTTTG	1720
	1721	AAAGTGCCAATGCTTTACATCTCACTCGGTAACATCGT	1760
	1761	GGGTGTTAGAAACTTTAGTGGGACTGCAGGAGTGATTATC	1800
10	1801	GACAGATTGAGTTCATTCAGTTACTGCAACACTCGAGG	1840
	1841	CTGAGTACAACCTTGAGAGAGGCCAGAAGGCTGTGAACGC	1880
15	1881	CCTCTTACCTCCACCAATCAGCTGGCTTGAAAACAAAC	1920
	1921	GTTACTGACTATCACATTGACCAAGTGTCCAACTTGGTCA	1960
	1961	CCTACCTTAGCGATGAGTTCTGCCTCGACGAGAACCGTGA	2000
20	2001	ACTCTCCGAGAAAGTTAACACGCCAAGCGTCTCAGCGAC	2040
	2041	GAGAGGAATCTCTGCAAGACTCCAACCTCAAAGACATCA	2080
	2081	ACAGGCAGCCAGAACGTGGTTGGGTGGAAGCACCGGGAT	2120
25	2121	CACCATCCAAGGAGGCGACGATGTGTTCAAGGAGAACTAC	2160
	2161	GTCACCCCTCTCCGGAACCTTCGACGAGTGCTACCCCTACCT	2200
30	2201	ACTTGTACCAGAAGATCGATGAGTCCAAACTCAAAGCCTT	2240

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2241	CACCAAGGTATCAACTTAGAGGCTACATCGAAGACAGCCAA	2280
5	2281 GACCTTGAAATCTACTCGATCAGGTACAATGCCAAGCACG	2320
	2321 AGACCGTGAATGTCCCAGGTACTGGTCCCTCTGGCCACT	2360
	2361 TTCTGCCAATCTCCCATTGGAAAGTGTGGAGAGCCTAAC	2400
10	2401 AGATGCGCTCCACACCTTGAGTGGAAATCCTGACTTGGACT	2440
	2441 GCTCCTGCAGGGATGGCGAGAAGTGTGCCACCATTCTCA	2480
15	2481 TCACTTCTCCTTGGACATCGATGTGGATGTACTGACCTG	2520
	2521 AATGAGGACCTCGGAGTCTGGTCATCTCAAGATCAAGA	2560
	2561 CCCAAGACGGACACGCAAGACTGGAACCTTGAGTTCT	2600
20	2601 CGAAGAGAAACCATTGGTCGGTGAAGCTCTCGCTCGTGTG	2640
	2641 AAGAGAGCAGAGAAGAAGTGGAGGGACAAACGTGAGAAC	2680
	2681 TCGAATGGAAACTAACATCGTTACAAGGAGGCCAAAGA	2720
25	2721 GTCCGTGGATGCTTGTTCGTGAACCTCCAATATGATCAG	2760
	2761 TTGCAAGCCGACACCAACATGCCATGATCCACGCCGCAG	2800
30	2801 ACAAACGTGTGCACAGCATTGAGGCTTACTGCCTGA	2840

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2841	GTTGTCCGTGATCCCTGGTGTGAACGCTGCCATCTTCGAG	2880
5	2881 GAACTTGAGGGACGTATCTTACCGCATTCTCCTTGTACG	2920
	2921 ATGCCAGAACGTCATCAAGAACGGTACTTACAACAATGG	2960
10	2961 CCTCAGCTGCTGGAATGTGAAAGGTACATGTGGACGTGGAG	3000
	3001 GAACAGAACAAATCAGCGTTCCGTCTGGTTGTGCCTGAGT	3040
	3041 GGGAAAGCTGAAGTGTCCCAAGAGGTTAGAGTCTGTCCAGG	3080
15	3081 TAGAGGCTACATTCTCCGTGTGACCGCTTACAAGGAGGGA	3120
	3121 TACGGTGAGGGTTGCGTGACCATCCACGAGATCGAGAACAA	3160
	3161 ACACCGACGAGCTTAAGTTCTCCAATGCGTCGAGGAAGA	3200
20	3201 AATCTATCCAACAAACACCGTTACTTGCAACGACTACACT	3240
	3241 GTGAATCAGGAAGAGTACGGAGGTGCCTACACTAGCCGTA	3280
25	3281 ACAGAGGTTACAACGAAGCTCCTCCGTTCCCTGCTGACTA	3320
	3321 TGCCTCCGTGTACGAGGAGAAATCCTACACAGATGGCAGA	3360
	3361 CGTGAGAACCTTGCAGTTCAACAGAGGTTACAGGGACT	3400
30	3401 ACACACCACCCAGTTGGCTATGTTACCAAGGAGCTTGA	3440

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3441 GTACTTCCTGAGACCGACAAAGTGTGGATCGAGATCGGT 3480

5 3481 GAAACCGAGGGAACCTTCATCGTGGACAGCGTGGAGCTTC 3520

3521 TCTTGATGGAGGAA 3534.

22. A structural gene of Claim 13 which encodes an  
10 insecticidal protein of *B.t.t.* having the sequence:

1 ATGACTGCAGACAACAACACCGAAGCCCTCGACAGTTCTA 40

41 CCACTAAGGATGTTATCCAGAAGGGTATCTCCGTTGTGGG 80

15 81 AGACCTCTGGCGTGGTTGGATTTCCCTTCGGTGGAGCC 120

121 CTCGTGAGCTTCTATACAAACTTCTAACACACCATTGGC 160

20 161 CAAGCGAGGACCCCTGGAAAGCATTCATGGAGCAAGTTGA 200

201 AGCTCTTATGGATCAGAAGATTGCAGATTATGCCAAGAAC 240

241 AAGGCTTGGCAGAACTCCAGGGCCTTCAGAACAAATGTGG 280

25 281 AGGACTACGTGAGTGCATTGTCCAGCTGGCAGAAGAACCC 320

321 TGTTAGCTCCAGAAATCCTCACAGCCAAGGTAGGATCAGA 360

361 GAGTTGTTCTCTCAAGCCGAATCCCACTTCAGAAATTCCA 400

30

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401	TGCCTAGCTTGCTATCTCCGGTTACGAGGTTCTTTCT	440
5	CACTACCTATGCTCAAGCTGCCAACACCCACTTGTTCTC	480
481	CTTAAGGACGCTCAAATCTATGGAGAAGAGTGGGGATACG	520
10	521 AGAAAGAGGACATTGCTGAGTTCTACAAGCGTCAACTAA	560
561	GCTCACCCAAGAGTACACTGACCATTGCGTGAATGGTAT	600
601	AACGTTGGTCTCGATAAGCTCAGAGGCTCTTCCTACGAGT	640
15	641 CTTGGGTGAACCTAACAGATAACAGGAGAGATGACCTT	680
681	GACTGTGCTCGATCTTATCGCACTCTTCCCTGTACGAT	720
721	GTGAGACTCTACCCAAAGGAAGTGAAAATGAGCTTACCA	760
20	761 GAGACGTGCTCACTGACCCTATTGTCGGAGTCAACAAACCT	800
801	TAGGGTTATGGAACCTACCTTCAGCAATATCGAAACTAC	840
25	841 ATTAGGAAACCACATCTCTCGACTATCTCACAGAATTCT	880
881	AATTCCACACAAAGGTTCAACCAGGATACTATGGTAACGA	920
921	CTCCTTCAACTATTGGTCCGGTAACTATGTTCCACCAGA	960
30	961 CCAAGCATTGGATCTAATGACATCATCACATCTCCCTCT	1000

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1001	ATGGTAACAAGTCCAGTGAACCTGTGCAGAACCTTGAGTT	1040	
5	CAACGGCGAGAAAGTCTATAGAGCCGTCGCAAACACCAAT	1080	
1081	CTCGCTGTGTGGCCATCCGCAGTTACTCAGGCGTCACAA	1120	
1121	AGGTGGAGTTAGTCAGTATAACGATCAGACCGATGAGGC	1160	
10	1161	CAGCACCCAGACTTACGACTCCAAACGTAACGTTGGCGCA	1200
1201	GTCTCTTGGGATTCTATCGACCAATTGCCTCCAGAAACCA	1240	
15	1241	CAGACGAACCATTGGAGAAGGGCTACAGCCACCAACTTAA	1280
1281	CTATGTGATGTGCTTCTTGATGCAAGGTTCCAGAGGGACC	1320	
1321	ATTCCAGTGTGACCTGGACACACACAAGTCCGTGGACTTCT	1360	
20	1361	TCAACATGATCGATAGCAAGAAGATCACTCAACTCCCTT	1400
1401	GGTGAAAGCCTACAAGCTGCAATCTGGTGCTTCCGTTGTC	1440	
25	1441	GCAGGTCCCAGATTCACTGGAGGTGACATCATCCAGTGCA	1480
1481	CAGAGAACGGCAGCGCAGCTACTATCTACGTGACACCTGA	1520	
1521	TGTGTCTTACTCTCAGAAGTACAGGGCACGTATTCAATTAC	1560	
30	1561	GCATCTACCAGCCAGATCACCTTCACACTCAGCTTGGATG	1600

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1601	GAGCACCCCTCAACCAGTATTACTTGTACAAGACCATCAA	1640
5	CAAAGGTGACACTCTCACATACAATAGCTCAACTTGGCA	1680
1681	AGTTTCAGCACACCATTGAACCTCTCAGGCAACAATCTTC	1720
10	1721 AGATCGGCGTCACCGGTCTCAGCGCCGGAGACAAAGTCTA	1760
1761	CATCGACAAGATTGAGTTCATCCCAGTGAAC	1791.

23. A structural gene of Claim 13 which encodes an  
insecticidal protein of *B.t. entomocidus* having the  
15 sequence:

1	ATGGAGGAGAACAAACCAAAACCAATGCATTCCATACAAC	40
41	GCTTGAGTAACCCAGAAGAGGTATTGCTGATGGAGAACG	80
20	CATTTCAACCGGTAACCTTCCATCGACATCTCCTTGTCC	120
81	TTGGTCCAGTTCTGGTCAGCAACTTCGTGCCAGGTGGTG	160
121	GGTTCCCTTGTGGACTAATTGACTTCGTCTGGGTATCGT	200
25	TGGTCCATCTCAATGGGATGCATTCCCTGGTGCAAATTGAG	240
161	CAGTTGATCAACGAGAGGATCGCTGAGTTGCCAGGAACG	280
201	CTGCCATCGCTAACCTGGAAGGATTGGGCAATAACTTCAA	320
241		
30		

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	321	CATCTATGTGGAGGCCTTCAAAGAGTGGGAAGAGGACCT	360
5	361	AACAACCCAGAGACCCGCACTAGGGTGATCGACAGATTCA	400
	401	GAATCTTGGACGGCCTCTTGGAGAGAGATATCCCATCCTT	440
	441	CAGAATCTCTGGCTTCGAAGTTCCCTCTTGTCCTGTAC	480
10			
	481	GCTCAAGCAGCTAATCTCACCTCGCTATCCTCGAGACA	520
	521	GTGTCATCTTGGGGAAAGGTGGGGATTGACCACTATCAA	560
15	561	CGTCAATGAGAATTACAACAGACTTATCAGGCACATTGAC	600
	601	GAGTACGCCGACCACTGTGCTAACACCTACAACCGTGGCT	640
	641	TGAACAATCTCCCTAAGTCTACTTATCAAGATTGGATTAC	680
20			
	681	CTACAACAGGTTGAGGAGAGACTTGACCCCTCACAGTTTG	720
	721	GACATTGCAGCTTCTTCCCGAACTATGACAACAGGAGAT	760
25	761	ACCCTATCCAACCAGTGGGTCAACTTACCAAGAGAAGTCTA	800
	801	TACTGACCCACTTATCAACTCAACCCCTCAGTTGCAAAGT	840
	841	GTCGCCAACCTCCCACATTCAACGTCACTGGAGTCCAGCC	880
30	881	GTATCAGGAACCCACACTTGTGACATCTTGAACAACCT	920

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921	TACTATCTCACCGATTGGTCAGCGTTGGCGTAACCTTC	960
5	TATTGGGGTGGACACAGGGTCATCTCCTCTCTTATTGGAG	1000
1001	GTGGGAACATTACCTCTCCTATCTATGGACGTGAGGCAA	1040
1041	CCAGGAGCCACCACGTAGTTCACCTCAACGGTCCAGTC	1080
10	1081 TTCAGAACCTTGTCTAACCTACCTTGAGATTGCTCCAGC	1120
1121	AACCTTGGCCAGCTCCACCTTCAACCTTAGAGGTGTTGA	1160
15	1161 GGGCGTTGAGTTCTACTCCTACCAACTCCTCACTTAC	1200
1201	AGAGGTAGAGGAACCGTTGATTCTTGACCGAACTCCCAC	1240
1241	CAGAGGACAATAGCGTGCCACCCAGGGAAAGGCTACTCCCA	1280
20	1281 CAGGTTGTGCCACGCAACCTCGTGCAGCGTTCCGGAAC	1320
1321	CCATTCTCACTACAGGAGTTGTGTTCTCATGGACTGATC	1360
1361	GTAGTGCTACTCTCACTAACCAATTGATCCCGAGAGGAT	1400
25	1401 CAATCAAATCCCATTGGTCAAGGGTTCCGTGTGGGGAA	1440
1441	GGAACTTCTGTACACAGGACCAGGCTTCACAGGAGGTG	1480
30	1481 ATATTCTAGAAGAACACTTTGGCGACTTGTGAGCCT	1520

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1521	CCAAGTTAACATCAACTCTCCAATTACTCAAAGATATCGT	1560
5	CTCAGGTTCGTTACGCATCTCCCGTGACGCTAGAGTCA	1600
1561	TCGTGCTCACCGGAGCAGCTCTACCGGTGTCGGTGGACA	1640
1601	AGTCTCCGTGAACATGCCACTCCAGAAGACTATGGAGATC	1680
10	GGCGAGAACTTGACATCCAGGACCTTCAGATAACCCGACT	1720
1641	TCTCTAACCCCTTCAGTTCCGTGCCAACCTGACATCAT	1760
1681	TGGCATTAGCGAACAAACCTCTCTTGAGCTGGTAGCATC	1800
15	TCATCTGGCGAATTGTACATTGACAAGATTGAGATCATTC	1840
1721	TTGCCGACGCTACCTCGAGGCTGAGTCTGACCTTGAGAG	1880
1761	AGCCCAGAAGGCTGTGAACGCCCTTTACCTCTTAAT	1920
20	CAGATTGGCTTGAAACTGACGTTACTGACTATCACATTG	1960
1801	ACCAAGTGTCCAACTTGGTCGACTGCCTTAGCGATGAGTT	2000
1841	CTGCCTCGACGAGAAGCGTGAACCTCCGAGAAAGTTAAA	2040
2041	CACGCCAAGCGTCTCAGCGACGAGAGGAATCTTGCAAG	2080
25	ACCCCAACTTCAGAGGCATAACAGGGCAGCCAGACCGTGG	2120
1881	2081	
1921		
1961		
2001		
2041		
30		

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2121	TTGGAGAGGAAGCACCGACATCACCATCCAAGGAGGCGAC	2160
5	5 2161 GATGTGTTCAAGGAGAACTACGTCACCCCTCCCAGGAAC	2200
2201	TGGACGAGTGCTACCCCTACCTACTTGTACCAGAAAGATCGA	2240
10	10 2241 TGAGTCCAAACTCAAAGCCTACACCAGGTATGAAC	2280
2281	2281 GGCTACATCGAAGACAGCCAAGACCTTGAAATCTACCTCA	2320
2321	2321 TCAGGTACAATGCCAAGCACCGAGATCGTGAATGTCCCAGG	2360
15	15 2361 TACTGGTCCCTCTGGCCACTTCTGCCAAATGCCATT	2400
2401	2401 GGGAAAGTGTGGAGAGCCTAACAGATGCGCTCCACACCTTG	2440
2441	2441 AGTGGAATCCTGACTTGGACTGCTCCTGCAGGGATGGCGA	2480
20	20 2481 GAAGTGTGCCACCATTCTCATCACTCACCTTGACATC	2520
2521	2521 2521 GATGTGGATGTACTGACCTGAATGAGGACCTCGGAGTCT	2560
25	25 2561 GGGTCATCTCAAGATCAAGACCCAAGACGGACACGCAAG	2600
2601	2601 2601 ACTTGGCAACCTTGAGTTCTGAAGAGAAACCATTGCTC	2640
2641	2641 2641 GGTGAAGCTCTCGCTCGTGTGAAGAGAGCAGAGAAGAAGT	2680
30	30 2681 GGAGGGACAAACGTGAGAAACTCCAAC	2720
	TCGAGACTAACAT	

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2721	CGTTTACAAGGAGGCCAAAGAGTCCGTGGATGCTTGTT	2760
5	GTGAACCTCCAATATGATAGGTTGCAAGTGGACACCAACA	2800
2801	TCGCCATGATCCACGCTGCAGACAAACGTGTGCACAGGAT	2840
10	TCGTGAGGCTTACTTGCCTGAGTTGTCGTGATCCCTGGT	2880
2881	GTGAACGCTGCCATCTTCGAGGAACTTGAGGGACGTATCT	2920
2921	TTACCGCATACTCCTTGTACGATGCCAGAAACGTCATCAA	2960
15	GAACGGTGACTTCAACAATGGCCTTTGTGCTGGAATGTG	3000
3001	AAAGGTCATGTGGACGTGGAGGAACAGAACAAATCACC GTT	3040
3041	CCGTCCCTGGTTATCCCTGAGTGGGAAGCTGAAGTGTCCCA	3080
20	AGAGGTTAGAGTCTGTCCAGGTAGAGGCTACATTCTCCGT	3120
3121	GTGACCGCTTACAAGGAGGGATACGGTGAGGGTTGCGTGA	3160
25	CCATCCACGAGATCGAGGACAACACCGACGAGCTTAAGTT	3200
3201	CTCCAAC TGCGTCGAGGAAGAAGTCTATCCAAACAACACC	3240
3241	GTTACTTGCAACAACTACACTGGGACCCAGGAAGAGTACG	3280
30	AAGGTACCTACACTAGCCGTAACCAAGGTTACGACGAAGC	3320

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3321	TTACGGAAACAATCCTCCGTTCTGCTGACTATGCCTCC	3360
5	GTGTACGAGGAGAAATCCTACACAGATGGCAGACGTGAGA	3400
3401	ACCCCTGCGAGTCCAACAGAGGTTACGGTGACTACACACC	3440
10	3441 ACTTCCAGCAGGCTATGTTACCAAGGACCTTGAGTACTTT	3480
3481	CCTGAGACCGACAAAGTGTGGATCGAGATCGGTGAAACCG	3520
3521	AGGGAACCTTCATCGTGGACAGCGTGGAGCTTCTCTTGAT	3560
15	3561 GGAGGAA 3567.	

24. A structural gene of Claim 13 which encodes a P2 insecticidal protein having the sequence:

20	1 ATGGACAAACAACGTCTTGAACCTCTGGTAGAACAAACCATCT	40
41	GCGACGCATAAACGTCGTGGCTCACGATCCATTAGCTT	80
81	CGAACACAAAGAGCCTCGACACTATTAGAAGGAGTGGATG	120
25	121 GAATGGAAACGTACTGACCACTCTCTACGTCGCACCTG	160
161	TGGTTGGAACAGTGTCCAGCTTCTCAAGAAGGAGTCGG	200
30	201 CTCTCTCATCGGAAAACGTATCTGTCCGAACCTGGGGT	240

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	241	ATCATTTCCATCTGGTCCACTAATCTCATGCAAGACA	280
5	281	TCTTGAGGGAGACCGAACAGTTCTCAACCAGCGTCTCAA	320
	321	CACTGATACTTGGCTAGAGTCAACGCTGAGTTGATCGGT	360
	361	CTCCAAGCAAACATTCTGTGAGTTCAACCAGCAAGTGGACA	400
10	401	ACTTCTTGAATCCAACCTCAGAACATCCTGTGCCCTTTCCAT	440
	441	CACTTCTTCCGTGAACACTATGCAGCAACTCTCCTCAAC	480
15	481	AGATTGCCTCAGTTTCAGATTCAAGGCTACCAGTTGCTCC	520
	521	TTCTTCCACTCTTGCTCAGGCTGCCAACATGCACTTGTC	560
	561	CTTCATACGTGACGTGATCCTCAACGCTGACGAATGGGA	600
20	601	ATCTCTGCAGCCACTCTTAGGACATACAGAGACTACTTGA	640
	641	GGAACATACACTCGTGATTACTCCAACATTGCATCAACAC	680
	681	TTATCAGACTGCCCTTCGTGGACTCAATACTAGGCTTCAC	720
25	721	GACATGCTTGAGTTCAAGGACCTACATGTTCTAACGTGT	760
	761	TTGAGTACGTCAGCATTGGAGTCTCTTCAAGTACCAGAG	800
30	801	CTTGATGGTGTCCCTCTGGAGCCAATCTCTACGCCCTGGC	840

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841	AGTGGACCACAGCAAACTCAGAGCTTCACAGCTCAGAACT	880
5	GGCCATTCTTGTATAGCTTGTCCAAGTCAACTCCAAC	920
921	CATTCTCAGTGGTATCTCTGGGACCAGACTCTCCATAACC	960
961	TTTCCCAACATTGGTGGACTTCCAGGCTCCACTACAACCC	1000
10	ATAGCCTTAACCTCTGCCAGAGTGAAC	1040
1001	TACAGACTACAGTGGAGGTGT	1080
1041	CAGCTCTGGATTGATTGGTGCAACTAACTTGAACCACAAAC	1120
15	TTCAATTGCTCCACCGTCTGCCACCTCTGAGCACACCGT	1160
1121	TTGTGAGGTCCCTGGCTTGACAGCGGTACTGATCGCGAAGG	1200
1161	AGTTGCTACCTCTACAAACTGGCAAACCGAGTCCTCCAA	1240
20	ACCACTCTAGCCTCGGTGTGGAGCTTCTGACGTG	1280
1201	ACCACTCTAGCCTCGGTGTGGAGCTTCTGACGTG	1320
1241	GGAATTCAAAC	1360
25	ACTTCCATTACAACCAGATTAGGAACATCGAGTCTC	1400
1281	CTCTGGTGTTCCTCTCGTCATCAGGAATGAAGACCTCACC	1440
1321	CGTCCACTTCATTACAACCAGATTAGGAACATCGAGTCTC	1480
1361	CATCCGGTACTCCAGGAGGTGCAAGAGCTTACCTCGTGTC	1520
30	TGTCCATAACAGGAAGAACACATCTACGCTGCCAACGAG	1560

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1441	AATGGCACCATGATTCACCTTGCACCAGAAGATTACACTG	1480
5	5 1481 GATTCACCATCTCTCCAATCCATGCTACCCAAGTGAACAA	1520
	1521 TCAGACACGCACCTTCATCTCCGAAAAGTTCGGAAATCAA	1560
	1561 10 GGTGACTCCTTGAGGTTCGAGCAATCCAACACTACCGCTA	1600
	1601 1601 GGTACACTTGAGAGGCAATGGAAACAGCTACAACCTTTA	1640
	1641 1641 CTTGAGAGTTAGCTCCATTGGTAACTCCACCATCCGTGTT	1680
15	1681 1681 ACCATCAACGGACGTGTTACACAGTCTCTAATGTGAACA	1720
	1721 1721 CTACAACGAACAATGATGGCGTTAACGACAACGGAGCCAG	1760
	1761 1761 ATTCAAGCGACATCAACATTGGCAACATCGTGGCCTCTGAC	1800
20	1801 1801 AACACTAACGTTACTTGGACATCAATGTGACCCCTCAATT	1840
	1841 1841 CTGGAACTCCATTGATCTCATGAACATCATGTTGTGCC	1880
25	1881 1881 AACTAACCTCCCTCCATTGTACTAA 1905.	

25. A plant transformation vector comprising a plant gene containing a structural gene of Claim 13.

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26. A structural gene sequence of Claim 13  
encoding a fusion protein comprising the N-terminal  
5 610 amino acids of *B.t.k.* HD-1 and the C-terminal 567  
amino acids of *B.t.k.* HD-73, said gene having the  
sequence:

1	ATGGACAAACAACCAAACATCAACGAATGCATTCCATACA	40
10	41 ACTGCTTGAGTAACCCAGAAGTTGAAGTACTTGGTGGAGA	80
	81 ACGCATTGAAACCGGTTACACTCCCATCGACATCTCCTTG	120
	121 TCCTTGACACAGTTCTGCTCAGCGAGTTCGTGCCAGGTG	160
15	.	.
	161 CTGGGTTCGTTCTCGGACTAGTTGACATCATCTGGGTAT	200
	201 CTTTGGTCCATCTCAATGGGATGCATTCTGGTGCAAATT	240
20	241 GAGCAGTTGATCAACCAACAGAGGGATCGAAGAGTTGCCAGGA	280
	281 ACCAGGCCATCTCTAGGTTGGAAGGATTGAGCAATCTCTA	320
	321 CCAAATCTATGCAGAGAGCTTCAGAGAGTGGGAAGCCGAT	360
25	361 CCTACTAACCCAGCTCTCCCGAGGAAATGCGTATTCAAT	400
	401 TCAACGACATGAACAGCGCCTTGACCACAGCTATCCCATT	440

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441	GTTCGCAGTCCAGAACTACCAAGTTCCCTCTCTGTCCGTG	480
5	5	5
481	TACGTTCAAGCAGCTAATCTCACCTCAGCGTGCTTCGAG	520
521	521	521
561	ACGTTAGCGTGTGGGCAAAGGTGGGGATTGATGCTGC	560
10	561	561
601	AACCATCAATAGCCGTTACAACGACCTTACTAGGCTGATT	600
641	601	601
641	GCTTGGAGCGTGTCTGGGTCTGATTCTAGAGATTGGAT	680
15	641	641
681	TAGATACAACCAGTTCAGGAGAGAATTGACCCTCACAGTT	720
721	681	681
721	TTGGACATTGTGTCTCTCTTCCGAACATATGACTCCAGAA	760
761	721	721
761	CCTACCCCTATCCGTACAGTGTCCCAACTTACCAAGAGAAAT	800
20	761	761
801	801	801
841	CTATACTAACCCAGTTCTTGAGAACTTCGACGGTAGCTTC	840
841	841	841
881	CGTGGTTCTGCCAAGGTATCGAAGGCTCCATCAGGAGCC	880
25	881	881
921	CACACTTGATGGACATCTTGAACAGCATAACTATCTACAC	920
921	921	921
961	CGATGCTCACAGAGGAGAGTATTACTGGTCTGGACACCAG	960
961	961	961
30	ATCATGGCCTCTCCAGTTGGATTCAAGCGGGCCCGAGTTA	1000
1001	1001	1001
1001	CCTTCCCTCTATGGAACATATGGAAACGCCGCTCCACA	1040

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1041	ACAAACGTATCGTTGCTCAACTAGGTCAAGGGTGTCTACAGA	1080
5	1081 ACCTTGTCTTCCACCTTGTACAGAAGACCCCTCAATATCG	1120
	1121 GTATCAACAAACCAAGCAACTTCCGTTCTTGACGGAACAGA	1160
	1161 GTTCGCCTATGGAACCTCTTCTAACTTGCCATCCGCTGTT	1200
10	1201 TACAGAAAGAGCGGAACCGTTGATTCCCTTGGACGAAATCC	1240
	1241 CACCACAGAACAAACAATGTGCCACCCAGGCAAGGATTCTC	1280
15	1281 CCACAGGTTGAGCCACGTGTCCATGTTCCGTTCCGGATTC	1320
	1321 AGCAACAGTTCCGTGAGCATCATCAGAGCTCCTATGTTCT	1360
	1361 CATGGATTCATCGTAGTGCTGAGTTCAACAATATCATTCC	1400
20	1401 TTCCCTCTCAAATCACCCAAATCCCATTGACCAAGTCTACT	1440
	1441 AACCTTGGATCTGGAACCTCTGTCGTGAAAGGACCAGGCT	1480
25	1481 TCACAGGAGGTGATATTCTTAGAAGAACCTCTCCTGGCCA	1520
	1521 GATTAGCACCCCTCAGAGTTAACATCACTGCACCACTTCT	1560
	1561 CAAAGATATCGTGTCAAGGATTGTTACGCATCTACCACTA	1600
30	1601 ACTTGCAATTCCACACCTCCATCGACGGAAGGCCTATCAA	1640

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	1641	TCAGGGTAACCTCTCCGCAACCATGTCAAGCGGCAGCAAC	1680
5	1681	TTGCAATCCGGCAGCTTCAGAACCGTCGGTTCACTACTC	1720
	1721	CTTTCAACTCTCTAACGGATCAAGCGTTTCACCCTAG	1760
	1761	CGCTCATGTGTTCAATTCTGGCAATGAAGTGTACATTGAC	1800
10	1801	CGTATTGAGTTGTGCCCTGCCGAAGTTACCCCTGAGGCTG	1840
	1841	AGTACAACCTTGAGAGAGGCCAGAAGGCTGTGAACGCCCT	1880
15	1881	CTTTACCTCCACCAATCAGCTTGGCTTGAAACTAACGTT	1920
	1921	ACTGACTATCACATTGACCAAGTGTCCAATTGGTCACCT	1960
	1961	ACCTTAGCGATGAGTTCTGCCTCGACGAGAAGCGTGAAC	2000
20	2001	CTCCGAGAAAGTTAACACACGCCAACGCGTCTCAGCGACGAG	2040
	2041	AGGAATCTCTTGCAGACTCCAATTCAAAGACATCAACA	2080
	2081	GGCAGCCAGAACGTGGTGGGTGGAAGCACCGGGATCAC	2120
25	2121	CATCCAAGGAGGCGACGATGTGTTCAAGGAGAACTACGTC	2160
	2161	ACCCCTCTCCGGAACCTTCGACGAGTGCTACCCCTACCTACT	2200
30	2201	TGTACCAGAAGATCGATGAGTCCAAACTCAAAGCCTTCAC	2240

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2241	CAGGTATCAACTTAGAGGCTACATCGAAGACAGCCAAGAC	2280
5	CTTGAAATCTACTCGATCAGGTACAATGCCAAGCACGAGA	2320
2321	CCGTGAATGTCCCAGGTACTGGTCCCTCTGGCCACTTC	2360
10	TGCCCAATCTCCCATTGGGAAGTGTGGAGAGCCTAACAGA	2400
2401	TGCGCTCCACACCTTGAGTGGAAATCCTGACTTGGACTGCT	2440
2441	CCTGCAGGGATGGCGAGAAGTGTGCCACCATTCTCATCA	2480
15	CTTCTCCTTGGACATCGATGTGGATGTACTGACCTGAAT	2520
2521	GAGGACCTCGGAGTCTGGTCATCTCAAGATCAAGACCC	2560
20	AAGACGGACACGCAAGACTTGGCAACCTTGAGTTCTCGA	2600
2601	AGAGAAACCATTGGTCGGTGAAGCTCTCGCTCGTGTGAAG	2640
2641	AGAGCAGAGAAGAAGTGGAGGGACAAACGTGAGAAACTCG	2680
25	AATGGGAAACTAACATCGTTACAAGGAGGCCAAAGAGTC	2720
2721	CGTGGATGCTTGTTCGTGAACTCCCAATATGATCAGTTG	2760
2761	CAAGCCGACACCAACATGCCATGATCCACGCCGCAGACA	2800
30	AACGTGTGCACAGCATTGTGAGGCTACTTGCCTGAGTT	2840

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2841	GTCCGTGATCCCTGGTGTGAACGCTGCCATCTCGAGGAA	2880	
5	CTTGAGGGACGTATCTTACCGCATTCTCCTTGTACGATG	2920	
2921	CCAGAAACGTCATCAAGAACGGTGACTTCAACAATGGCCT	2960	
2961	CAGCTGCTGGAATGTGAAAGGTATGTGGACGTGGAGGAA	3000	
10	3001	CAGAACAAATCAGCGTTCCGTCCTGGTTGTGCCTGAGTGGG	3040
3041	AAGCTGAAGTGTCCAAGAGGTTAGAGTCTGTCCAGGTAG	3080	
15	3081	AGGCTACATTCTCCGTGTGACCGCTTACAAGGAGGGATAC	3120
3121	GGTGAGGGTTGCGTGACCATCCACGAGATCGAGAACACA	3160	
3161	CCGACGAGCTTAAGTTCTCAACTGCGTCGAGGAAGAAAT	3200	
20	3201	CTATCCAAACAACACCGTTACTTGCAACGACTACACTGTG	3240
3241	AATCAGGAAGAGTACGGAGGTGCCTACACTAGCCGTAACA	3280	
3281	GAGGTTACAACGAAGCTCCTCCGTTCTGCTGACTATGC	3320	
25	3321	CTCCGTGTACGAGGGAGAAATCCTACACAGATGGCAGACGT	3360
3361	GAGAACCCCTGCGAGTTAACAGAGGTTACAGGGACTACA	3400	
30	3401	CACCACTTCCAGTTGGCTATGTTACCAAGGAGCTTGAGTA	3440

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3441 CTTTCCTGAGACCGACAAAGTGTGGATCGAGATCGGTGAA 3480  
5 3481 ACCGAGGGAACCTTCATCGTGGACAGCGTGGAGCTCTCT 3520  
3521 TGATGGAGGAA 3531.

27. A method of Claim 4 further comprising removal  
10 of sequences comprising more than five consecutive A+T  
or G+C bases.

28. A structural gene sequence of Claim 13  
comprising a majority of plant preferred codons.

29. A structural gene encoding the coat protein of  
15 potato leaf roll virus, said gene having the sequence:

1 ATGAGTACTGTCGTGGTTAAGGGAAACGTGAACGGTGGTG 40  
41 TTCAACAAACCTAGAAGGAGAAGAAGGCAATCCCTTCGTAG 80  
20 81 GAGAGCTAACAGAGTTCAGCCAGTGGTTATGGTCACTGCT 120  
121 CCTGGGCAACCAAGAAGGAGAAGAAGGAGAAGAGGAGGTA 160  
25 161 ATCGCAGATCAAGAAGAAGACTGGAGTTCCCAGAGGAAGAGG 200  
201 TTCAAGCGAGACATTCTGTGTTACAAAGGACAACCTCGTG 240  
241 GGCAACTCCCAAGGAAGTTCACCTTCGGACCAAGTGT 280  
30 281 CAGACTGTCCAGCATTCAAGGATGGAATACTCAAGGCTTA 320

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	321	CCATGAGTACAAGATCACAAGTATCTGCTTCAGTCGTC	360
5	361	AGCGAGGCCTTTCCACCTCTCCAGGCTCCATCGCTTATG	400
	401	AGTTAGATCCACATTGCAAAGTTCATCCCTCCAGTCCTA	440
	441	CGTCAACAAGTTCAAATCACAAAGGGTGGTGCTAAGACC	480
10			
	481	TATCAAGCTCGTATGATCAACGGAGTTGAATGGCACGATT	520
	521	CTTCTGAGGATCAGTGCAGAACCTTGGAAAGGAAATGG	560
	561	AAAGTCTTCAGATCCAGCTGGATCTTCAGAGTTACCATC	600
15			
	601	AGAGTTGCTTTCAAAACCCAAAG	624.

30. A chimeric plant gene which comprises a structural coding sequence encoding an insecticidal protein of *Bacillus thuringiensis*, said structural coding sequence being modified to reduce the number of putative polyadenylation signals within said structural coding sequence.

31. A chimeric plant gene of Claim 30 in which the polyadenylation signals are selected from the group consisting of AATAAA, AATAAT, AACCAA, ATATAA, AATCAA, ATACTA, ATAAAA, ATGAAA, AAGCAT, ATTAAT, ATACAT, AAAATA, ATTAAA, AATTAA, AATACA and CATAAA.

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32. A chimeric plant gene of Claim 31 in which  
said structural coding sequence is further modified to  
reduce the number of ATTAA sequences within said  
5 structural coding sequence.

33. A chimeric plant gene of Claim 32 in which  
said structural coding sequence is substantially  
devoid of polyadenylation signals and ATTAA sequences.

34. A transformed plant cell containing a gene of  
10 Claim 33.

35. A transformed plant cell of Claim 34 selected  
from the group consisting of soybean, cotton, alfalfa,  
oilseed rape, flax, tomato, sugarbeet, sunflower,  
potato, tobacco, maize, rice and wheat.

15 36. A plant comprising transformed plant cells of  
Claim 34.

37. A plant of Claim 36 which comprises plant  
cells of Claim 35.

38. A seed produced by a plant of Claim 36.

20

25

30

## DETERMINATION OF DNA REGIONS IN GENES TO MODIFY BY SITE-DIRECTED MUTAGENESIS FOR INCREASED EXPRESSION IN PLANTS

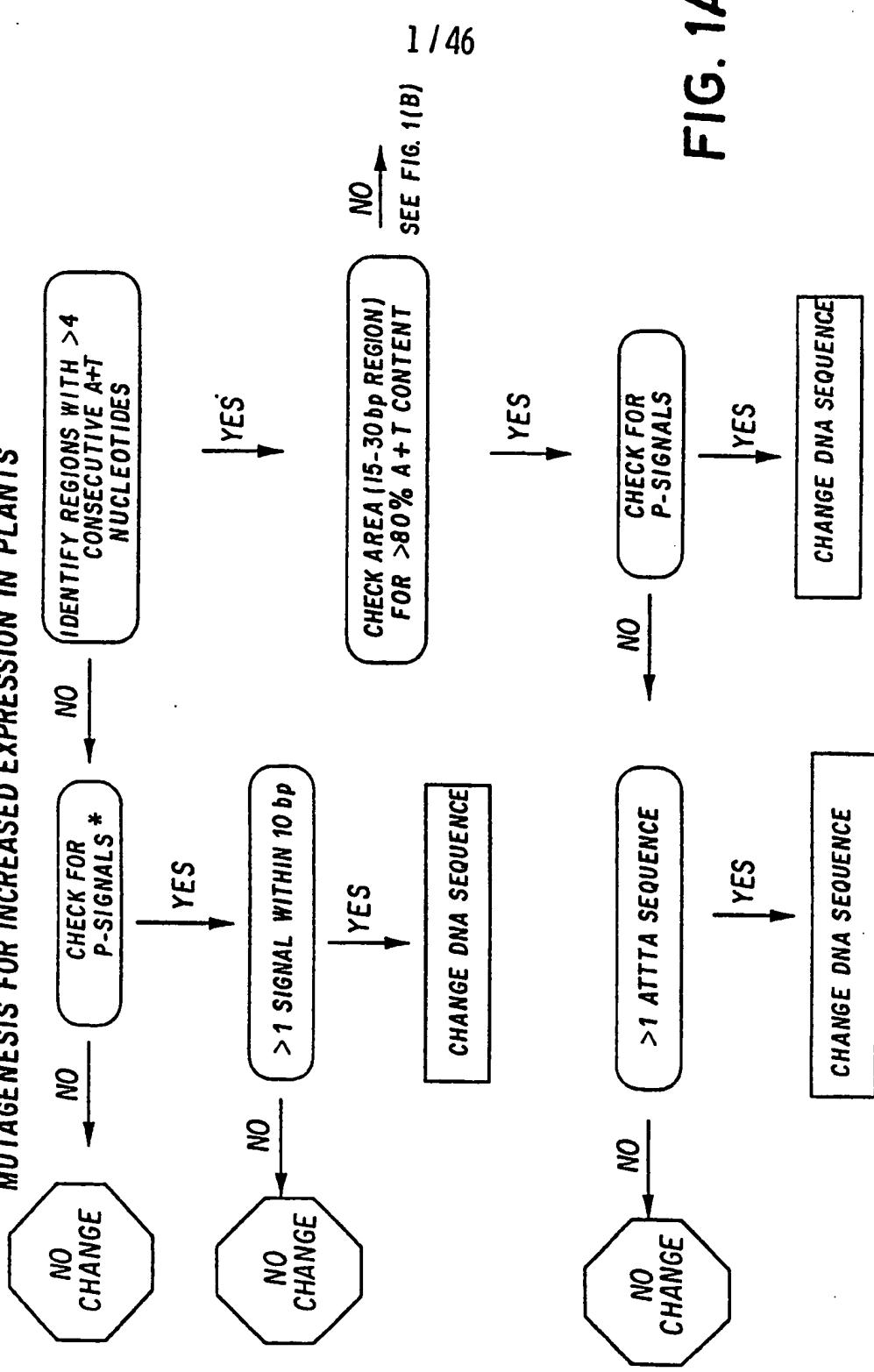


FIG. 1A

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SEE FIG. 1(B)

**CHECK AREA (13-300P REGION)  
FOR >80% A+T CONTENT**

**CHECK FOR  
P-SIGNALS**

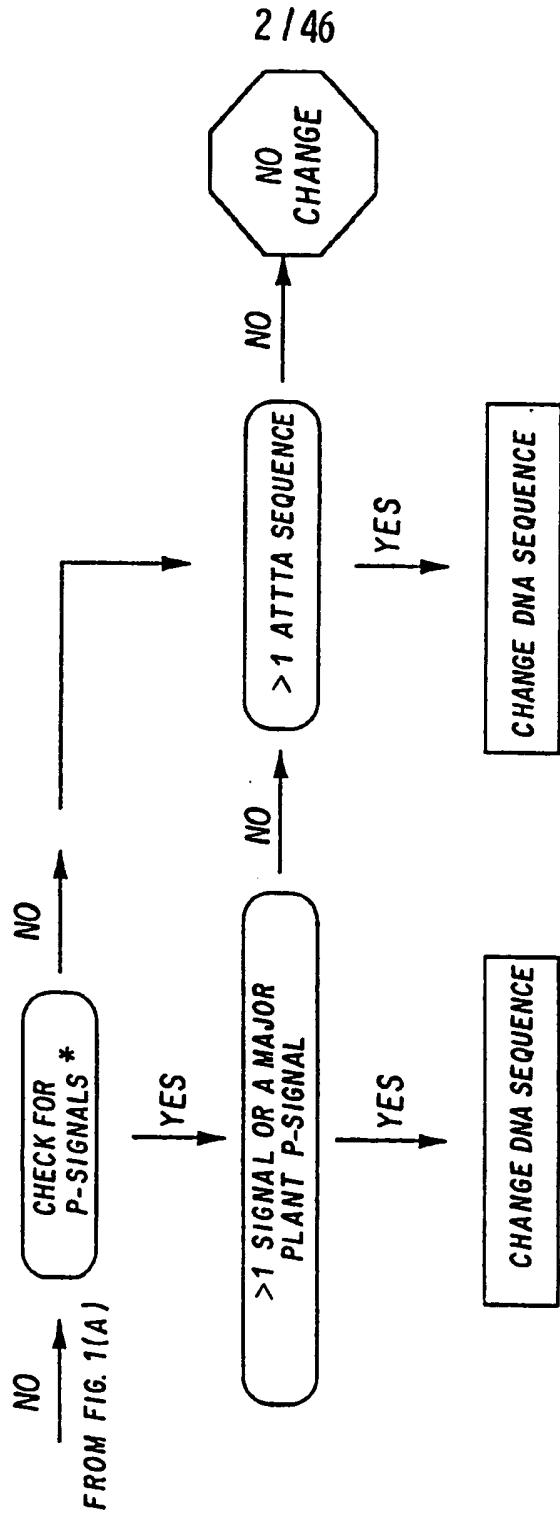
NO

## CHANGE DNA SEQUENCE

## CHANGE DNA SEQUENCE

\* POLYADENYLATION SIGNAL SEQUENCES

**DETERMINATION OF DNA REGIONS IN GENES TO MODIFY BY SITE-DIRECTED  
MUTAGENESIS FOR INCREASED EXPRESSION IN PLANTS**



\* POLYADENYLATION SIGNAL SEQUENCES

**FIG. 1B**

**SUBSTITUTE SHEET**

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1	ATGGCTATAGAAACTGGTTACACCCCAATCGATATTCCT	40
41	TGTCGCTAACGCAATTCTTTGAGTGAATTGTTCCGG	80
81	TGCTGGATTGTGTTAGGACTAGTTGATATAATATGGGA	120
	T C	
121	ATTTTGTCCTCTCAATGGGACGCATTCTGTACAAA	160
161	TTGAACAGTTAACCAAGAACATAGAAGAACATCGCTAG	200
	C C C G C G	
201	GAACCAAGCCATTCTAGATTAGAAGGACTAACGAACTT	240
	T	
241	TATCAAATTACGCAGAACATCTTTAGAGAGTGGGAAGCAG	280
281	ATCCTACTAATCCAGCATTAAAGAGAACAGATGCGTATTCA	320
321	ATTCAATGACATGAACAGTGCCCTACAACCGCTATTCT	360
361	CTTTTGCAGTTCAAAATTATCAAGTTCTCTTTATCAG	400
	CC C C	
401	TATATGTTCAAGCTGCAAATTACATTATCAGTTTGAG	440
	G C C C C C	
441	AGATGTTCAAGTGGACAAAGGTGGGGATTGATGCC	480
481	GCGACTATCAATAGTCGTTATAATGATTTAACTAGGCTA	520
521	TTGGCAACTATACAGATCATGCTGTACGCTGGTACAATAC	560
561	GGGATTAGAGCGTGTATGGGGACCGGATTCTAGAGATTGG	600
601	ATAAGATATAATCAATTAGAACAGAACACTAACACTG	640
	C G C C G C G T	
641	TATTAGATATCGTTCTCTATTCCGAACTATGATAGTAG	680
681	AACGTATCCAATTGAAACAGTTCCCAATTAAACAAGAGAA	720

FIG. 2A

SUBSTITUTE SHEET

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721	ATTTATACAAACCCAGTATTAGAAAATTTGATGGTAGTT	760
761	TTCGAGGCTCGGCTCAGGGCATAGAAGGAAGTATTAGGAG	800
801	TCCACATTTGATGGATATACTTAATAGTATAACCATCTAT	840
841	ACGGATGCTCATAGAGGAGAATATTATTGGTCAGGGCATC C C C T C	880
881	AAATAATGGCTTCTCCTGTAGGGTTTCGGGCCAGAATT G C	920
921	CACTTTCCGCTATATGGAACATGGAAATGCAGCTCCA	960
961	CAACAAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATA	1000
1001	GAACATTATCGTCCACCTTATATAGAAGACCTTTAATAT C	1040
1041	AGGGATAAATAATCAACAACTATCTGTTCTGACGGGACA C C C C	1080
1081	GAATTGCTTATGGAACCTCCTCAAATTGCCATCCGCTG	1120
1121	TATACAGAAAAAGCGGAACGGTAGATCGCTGGATGAAAT	1160
1161	ACCGCCACAGAATAACAACGTGCCACCTAGGCAAGGATT	1200
1201	AGTCATCGATTAAGCCATGTTCAATGTTCGTCAGGCT	1240
1241	TTAGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTT	1280
1281	CTCTGGATACATCGTAGTGCTGAATTAAATAATATAATT G C C C C C	1320
1321	CCTTCATCACAAATTACACAAATACCTTAAACAAAATCTA C C C ACC C G	1360
1361	CTAATCTTGGCTCTGGAACCTCTGTCGTTAAAGGACCAGG	1400

FIG. 2B

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1401	ATTTACAGGAGGAGATATTCTCGAAGAACCTCACCTGGC	1440
1441	CAGATTCAACCTTAAGAGTAAATATTACTGCACCATTAT	1480
1481	CACAAAGATATCGGGTAAGAATTGCTACGCTTCTACCAC	1520
1521	AAATTTACAATTCCATACATCAATTGACGGAAGACCTATT CC T G C	1560
1561	AATCAGGGGAATTTTCAGCAACTATGAGTAGTGGAGTA	1600
1601	ATTTACAGTCCGGAAGCTTTAGGACTGTAGGTTTACTAC	1640
1641	TCCGTTAACCTTCAAATGGATCAAGTGTATTTACGTTA	1680
1681	AGTGCTCATGTCTCAATTCAAGGCAATGAAGTTATATAG	1720
1721	ATCGAATTGAATTGTTCCGGCA	1743

**FIG. 2C****SUBSTITUTE SHEET**

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1	ATGGATAACAATCCGAACATCAATGAATGCATTCTTATA	40
	C C A C C A	
41	ATTGTTAACGAACTGGTTACACCCCAATCGATATTCCTTG	80
	C C G A T C T	
81	AAGAATAGAAACTGGTTACACCCCAATCGATATTCCTTG	120
	C C T C T C C C	
121	TCGCTAACGCAATTCTTTGAGTGAATTGTTCCCGGTG	160
	CT G A G G C C C G C G A	
161	CTGGATTTGTGTTAGGACTAGTTGATATAATATGGGAAT	200
	G C T C C C C C T	
201	TTTGCGTCCCTCTCAATGGGACGCATTCTTGTACAAATT	240
	C A T C G G G	
241	GAACAGTTAACCAAAAGAATAGAAGAATTGCTAGGA	280
	G G C G G C G C	
281	ACCAAGCCATTCTAGATTAGAAGGACTAACGAAATCTTA	320
	G C G G T G C	
321	TCAAATTTACGCAGAACATCTTTAGAGAGTGGGAAGCAGAT	360
	C C T GAGC C C	
361	CCTACTAATCCAGCATTAAAGAGAAGAGATGCGTATTCAAT	400
	C T C C C G A	
401	TCAATGACATGAACAGTGCCCTTACAACCGCTATTCTCT	440
	C C T G C A C AT	
441	TTTGCAGTTCAAAATTATCAAGTTCTCTTTATCAGTA	480
	G C C G C C C G G C G	
481	TATGTTCAAGCTGCAAATTACATTATCAGTTTGAGAG	520
	C A T C T C C C A G C G C T C	
521	ATGTTCAAGCTGCAAATTACATTATCAGTTTGAGAG	560
	C A G C G C T	
561	GAATCAATAGCTGTTATAATGATTTAACCTAGGCTTATT	600
	A C C C C C C T G	
601	GGCAACTATACAGATCATGCTGTaCGCTGGTACAATACGG	640
	A C C C C T T C T	
641	GATTAGAGCGTGTATGGGGACCGGATTCTAGAGATTGGAT	680
	C G C T T C T	

FIG. 3A

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681	AAGATATAATCAATTAGAAGAGAATTAAACACTAACTGTA T C C G C G G C C A T	720
721	TTAGATATCGTTCTCTATTCCGAACATATGATAGTAGAA G C T G C C C T C C	760
761	CGTATCCAATTGAAACAGTTCCCAATTAAACAAGAGAAAT C C T C T G C T C	800
801	TTATACAAACCCAGTATTAGAAAATTGATGGTAGTTT C T T C T G C C C C	840
841	CGAGGCTCGGCTCAGGGCATAGAAGGAAGTATTAGGAGTC T T T C A T C C T C C	880
881	CACATTTGATGGATATACTTAATAGTATAACCATCTATAC C C T G C C T C	920
921	GGATGCTCATAGAGGAGAATATTATTGGTCAGGGCATCAA C C G C T A C G	960
961	ATAATGGCTCTCCTGTAGGGTTTCGGGCCAGAATTCA C C A T A C A G C C G T	1000
1001	CTTTCCGCTATATGGAACTATGGAAATGCAGCTCCACA C T C C C	1040
1041	ACAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATAGA C T C C	1080
1081	ACATTATCGTCCACCTTATATAGAAGACCTTTAATATAG C G T G C C C	1120
1121	GGATAAATAATCAACAACTATCTGTTCTGACGGGACAGA T C C C G T C A	1160
1161	ATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTGTA G C C T T C T	1200
1201	TACAGAAAAGCGGAACGGTAGATTGCTGGATGAAATAC G C T C T C C	1240
1241	CGCCACAGAATAACAACGTGCCACCTAGGCAAGGATTAG A C T C CTC	1280
1281	TCATCGATTAAGCCATGTTCAATGTTCGTCAGGCTTT C C A G G C G C C A C	1320
1321	AGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTTCT C C T C C G C C C	1360
1361	CTTGGATACATCGTAGTGCTGAATTAAATAATTCC A T G C C C	1400

**FIG. 3B**  
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1401	TTCATCACAAATTACACAAATACCTTTAACAAAATCTACT	1440
	C T C C C A G C G	
1441	AATCTTGGCTCTGGAACTTCTGTGCGTTAACAGGACCGAGGAT	1480
	C A G C	
1481	TTACAGGAGGAGATATTCTCGAAGAACCTCACCTGGCCA	1520
	C T A T	
1521	GATTTCAACCTTAAGAGTAAATATTACTGCACCATTATCA	1560
	AGC C C T C C C T T	
1561	CAAAGATATCGGGTAAGAATTGCTACGCTTCTACCACAA	1600
	T C G T A A	
1601	ATTTACAATTCCATACATCAATTGACGGAAGACCTATTAA	1640
	C G C C C G C	
1641	TCAGGGGAATTTTCAGCAACTATGAGTAGTAGGGAGTAAT	1680
	T C C C C T C A C C C C	
1681	TTACAGTCCGGAAGCTTTAGGACTGTAGGTTTACTACTC	1720
	G A C C A C C C C	
1721	CGTTTAACCTTCAAATGGATCAAGTGTATTACGTTAACG	1760
	T C C T C C T C C T	
1761	TGCTCATGTCTCAATTCAAGGCAATGAAGTTATATAGAT	1800
	C G T G C T C	
1801	CGAATTGAATTGTTCCGGCAGAACGTAACCTTGAGGCAG	1840
	T G G T C T C C T	
1841	AATAT 1845	
	G C	

FIG. 3C

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1	ATGGATAACAATCCGAACATCAATGAATGCATTCTTATA	40
	C C A C A C	
41	ATTGTTAACGTAACCTGAAAGTAGAAGTATTAGGTGGAGA	80
	C C G A T C T	
81	AAGAATAGAAACTGGTTACACCCCAATCGATATTCCTG	120
	C C T C T C C C	
121	TCGCTAACGCAATTCTTTGAGTGAATTGTCTCCGGTG	160
	CT G A G G C C C G C G A	
161	CTGGATTGTGTTAGGACTAGTTGATATAATATGGGAAT	200
	G C T C C C C C T	
201	TTTGGTCCCTCTCAATGGGACGCATTCTTGTACAAATT	240
	C A T C G G G	
241	GAACAGTTAACCAACAAAGAATAGAAGAATTGCTAGGA	280
	G G C G G C G C	
281	ACCAAGCCATTCTAGATTAGAAGGACTAAGCAATCTTA	320
	G C G G T G C	
321	TCAAATTACGCAGAACATCTTTAGAGAGTGGGAAGCAGAT	360
	C C T GAGC C C	
361	CCTACTAACCCAGCATTAAAGAGAAGAGATGCGTATTCAAT	400
	C T C C C G A	
401	TCAATGACATGAACAGTGCCTTACAACCGCTATTCTCT	440
	C C T G C A C AT	
441	TTTGCAAGTTCAAAATTATCAAGTTCTCTTTATCAGTA	480
	G C C G C C C G G	
481	TATGTTCAAGCTGCAAATTACATTATCAGTTTGAGAG	520
	C A T C T C C C A G C G C T C	
521	ATGTTCAAGTGTGGACAAAGGTGGGATTGATGCCGC	560
	C A G C G C T	
561	GACTATCAATAGTCGTTATAATGATTAACTAGGGCTTATT	600
	A C C C C C C T G	
601	GGCAACTATACAGATTATGCTGTACGCTGGTACAATACGG	640
	A C C C C C T T C T	
641	GATTAGAACGTGTATGGGGACCGGATTCTAGAGATTGGGT	680
	C G G C T T A	

FIG. 4A

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681	AAGGTATAATCAATTAGAAGAGAATTAACACTAACTGTA T A C C G C G G C C A T	720
721	TTAGATATCGTTGCTCTGTTCCCGAATTATGATAGTAGAA G C T G T C C C T C C	760
761	GATATCCAATTGAAACAGTTCCCAATTAAACAAGAGAAAT CC C T C T G C T C C	800
801	TTATACAAACCCAGTATTAGAAAATTTGATGGTAGTTT C T T C T G C C C C C C	840
841	CGAGGGCTCGGCTCAGGGCATAGAAAGAAGTATTAGGAGTC T T T C A T C G C T C C	880
881	CACATTGATGGATATACTAACAGTATAACCACATCTATAC C C T G C T C C	920
921	GGATGCTCATAGGGTTATTATTATTGGTCAGGGCATCAA C A A G G C T A C G	960
961	ATAATGGCTTCTCCTGTAGGGTTTCGGGGCCAGAATTCA C C A T A C A G C C G T	1000
1001	CTTTCCGCTATATGGAACATATGGAAATGCAGCTCCACA C T C C C	1040
1041	ACAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATAGA C T C C	1080
1081	ACATTATCGTCCACTTATATAGAAGACCTTTAATATAG C G T C G C C C C	1120
1121	GGATAAATAATCAACAACTATCTGTTCTGACGGGACAGA T C C C G T C A	1160
1161	ATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTGTA G C C T T C T	1200
1201	TACAGAAAAAGCGGAACGGTAGATTGCTGGATGAAATAC G C T C T C C C	1240
1241	CGCCACAGAATAACAACGTGCCACCTAGGCAAGGATTAG A C T C C T C	1280
1281	TCATCGATTAAGCCATGTTCAATGTTCGTTCAAGGCTTT C C A G G C G C C C A C	1320
1321	AGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTTCT C C T C C G C C C	1360
1361	CTTGGATACATCGTAGTGCTGAATTAAATAATTGC C G C C C C C	1400

**FIG. 4B**  
**SUBSTITUTE SHEET**

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1401	ATCGGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC	1440
	C	
1441	TTTCTTTAATGGTCTGTAATTCAGGACCAGGATTAC	1480
	C C C C	C
1481	CTGGTGGGGACTTAGTTAGATTAAATAGTAGTGGAAATAA	1520
	A C C C C C	
1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
1561	CCATCGACATCTACCAGATATCGAGTTCGTGTACGGTATG	1600
	C A GA	
1601	CTTCTGTAACCCGATTCAACCTAACGTTAATTGGGTAA	1640
	G T	
1641	TTCATCCATTTCACAGTACAGTACAGCTACAGCTACG	1680
	C C T C	
1681	TCATTAGATAATCTACAATCAAGTGATTTGGTTATTTG	1720
	C G C C C C	
1721	AAAGTGCCAATGCTTTACATCTTCATTAGGTAAATAGT	1760
	C C C C	
1761	AGGTGTTAGAAATTTAGTGGACTGCAGGAGTGATAATA	1800
	G C T C	
1801	GACAGATTTGAATTTATTCCAGTTACTGCAACACTCGAGG	1840
	C G C	
1841	CTGAATATAATCTGGAAAGAGCGCAGAAGGCGGTGAATGC	1880
	A TGCG	
1881	GCTGTTACGTCTACAAACCAACTAGGGCTAAAACAAAT	1920
	CTGT ACGTCTACA C AGCT G ACTC G CA TG	
1921	G 1921	

FIG. 4C

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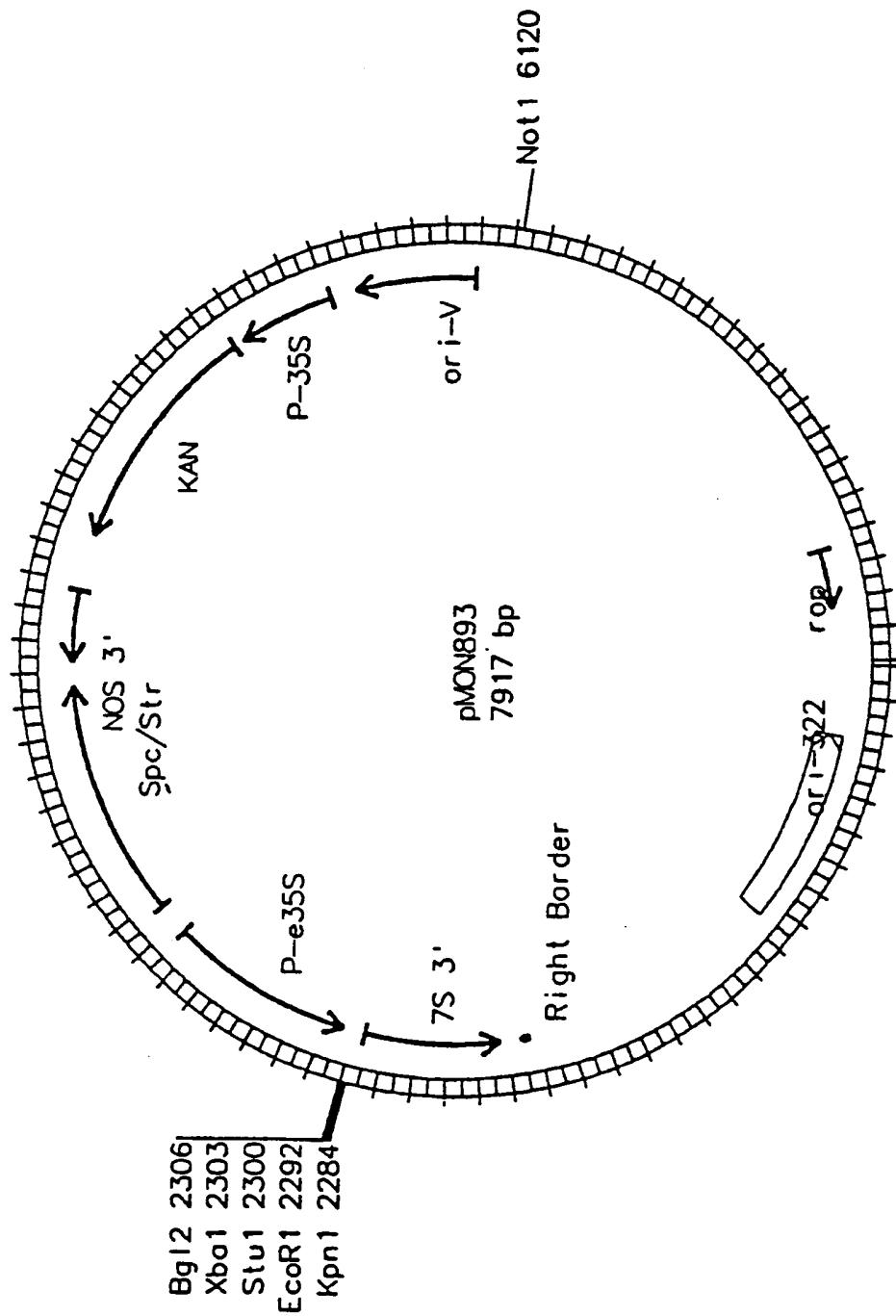


FIG. 5

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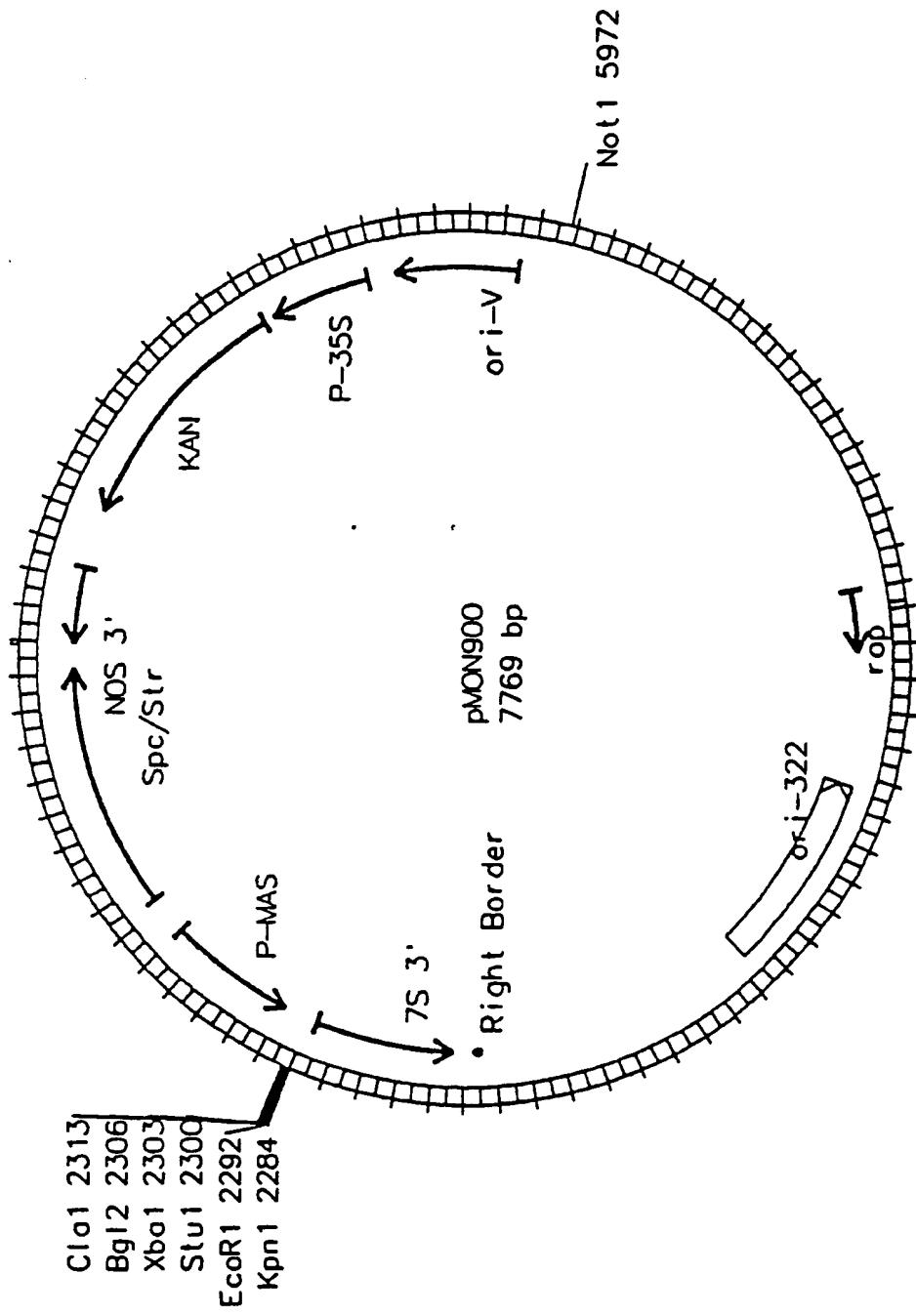


FIG. 6

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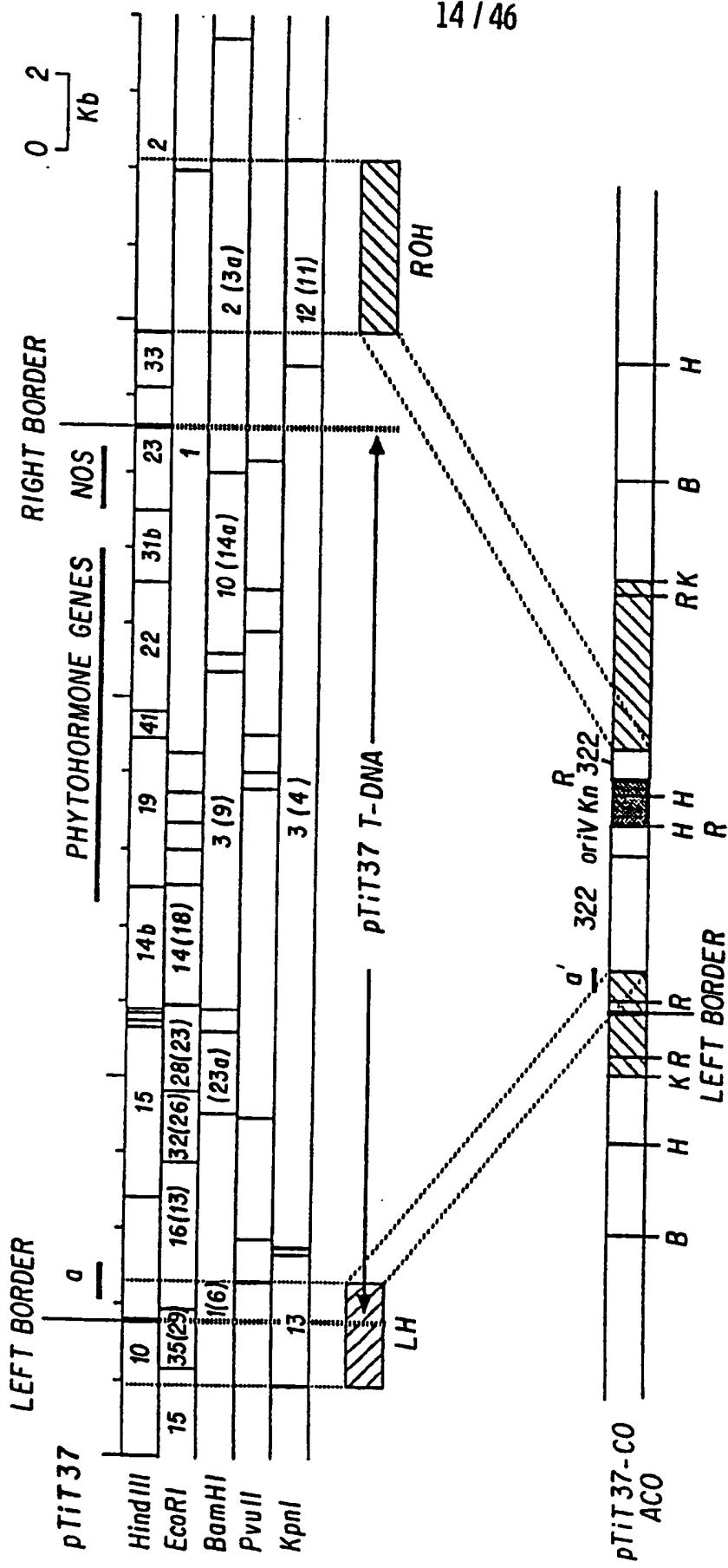


FIG. 7

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1	GAAAGAATAGAAACTGGTTACACCCCAATCGATATTCCT ATGGCC T C T C C C	40
41	TGTCGCTAACGCAATTCTTTGAGTGAATTGTTCCCGG CT G A G G C C C G C G A	80
81	TGCTGGATTGTGTTAGGACTAGTTGATATAATATGGGG G C T C C C C C T	120
121	ATTTTGGTCCCTCTCAATGGGACGCATTCTGTACAAA C A T C G G	160
161	TTGAACAGTTAACCAACAAAGAATAGAAGAAATTGCTAG G G C G G C G C	200
201	GAACCAAGCCATTCTAGATTAGAAGGACTAAGCAATCTT G C G G T G C	240
241	TATCAAATTACGCAGAACATCTTTAGAGAGTGGGAAGCAG C C T GAGC C C	280
281	ATCCTACTAATCCAGCATTAAAGAGAAAGAGATGCGTATTCA C TC C C G A	320
321	ATTCAATGACATGAACAGTGCCTTACAACCGCTATTCTT C C T G C A C A	360
361	CTTTTGCAAGTTCAAAATTATCAAGTTCTCTTTATCAG T G C C G C C C G C	400
401	TATATGTTCAAGCTGCAAATTACATTATCAGTTGAG G C A T C T C C C A G C G C T C	440
441	AGATGTTCAAGCTGCAAATTACATTATCAGTTGAG C AGC G C T	480
481	GCGACTATCAATAGTCGTTATAATGATTTAACTAGGCTTA A C C C C C T G	520
521	TTGGCAACTATACAGATTATGCTGTACGCTGGTACAATAC A C C C C T T C	560
561	GGGATTAGAACGTGTATGGGGACCGGGATTCTAGAGATTGG T C G G C T T	600
601	GTAAGGTATAATCAATTAGAAGAGAATTAAACACTAAG A T A C C G C G G C C A	640
641	TATTAGATATCGTTGCTCTGTTCCCGAATTATGATAGTAG T G C T G T C C C T C	680

FIG.8A

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681	AAGATATCCAATTGAAACAGTTCCCAATTAAACAAGAGAA	720
	CC C T C T G C T C	
721	ATTTATACAAACCCAGTATTAGAAAATTTGATGGTAGTT	760
	C T TC T G C C C C	
761	TTCGAGGCTCGGCTCAGGGCATAGAAAGAAGTATTAGGAG	800
	C T T T C A T C G CTCC C	
801	TCCACATTTGATGGATATACTAACAGTATAACCATCTAT	840
	C C C CT G C T C	
841	ACGGATGCTCATAGGGTTATTATTATTGGTCAGGGCATC	880
	C C A AG G C T A C	
881	AAATAATGGCTTCTCCTGTAGGGTTTCGGGCCAGAATT	920
	G C C A T A CAGC C G	
921	CACTTTCCGCTATATGGAACTATGGGAAATGCAGCTCCA	960
	T C T C C C	
961	CAACAAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATA	1000
	C T C C	
1001	GAACATTATCGTCCACTTTATATAGAACGACCTTTAATAT	1040
	C G T C G C C C	
1041	AGGGATAAATAATCAACAACTATCTGTTCTGACGGGACA	1080
	C T C C C G T C A	
1081	GAATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTG	1120
	G C C T T C	
1121	TATACAGAAAAAGCGGAACGGTAGATTCGCTGGATGAAAT	1160
	T G C T C T C	
1161	ACCGCCACAGAATAACAACGTGCCACCTAGGCAAGGATT	1200
	C A C T C C	
1201	AGTCATCGATTAAGCCATGTTCAATGTTCGTTCAAGGCT	1240
	TCC CA G G C G C C A	
1241	TTAGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTT	1280
	C C C TCC G C C C	
1281	CTCTGGATACATCGTAGTGCTGAATTAAATAATTAAATT	1320
	C G C C C C C	
1321	GCATCGGATAGTATTACTCAAATCCCTGCAGTGAAGGGAA	1360
	C	
1361	ACTTTCTTTAATGGTTCTGTAATTCAAGGACCAGGATT	1400
	C C C C	

FIG.8B

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1401	TA C	CTGGTGGGACTTAGTTAGATTAAATAGTAGTGGAAAT	1440
	A	CC	CC C C
1441	AACATT	CAGAATAGAGGGTATATTGAAGTTCCAATTCACT	1480
1481	TCCC	CATCGACATCTACCAGATATCGAGTTCGTGTACGGTA	1520
	C	A	GA
1521	TGCTT	CTGTAAACCCGATTCACCTAACGTTAATTGGGGT	1560
	G	T	
1561	AATT	CATCCATTTCACAGTACAGCTACAGCTA	1600
	C C	T	
1601	CGTC	CATTAGATAATCTACAAATCAAGTGATTTGGTTATTT	1640
	C C G	C C C C	C
1641	TGAAAGT	GCCAATGCTTTACATCTTCATTAGGTAATATA	1680
	C C	C C	
1681	GTAGGT	GTTAGAAATTAGGGACTGCAGGAGTGATAA	1720
	G	C	T
1721	TAGACAGATT	GAATTATTCCAGTTACTGCAACACTCGA	1760
	C	C G C	
1761	GGCTGAA	1767	
		G	

FIG. 8C

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1	ATGGATAACAATCCGAACATCAATGAATGCATTCTTATA	40
	C C A C A C	
41	ATTGTTAACGAAACCTGAAGTAGAAGTATTAGGTGGAGA	80
	C C G A T C T	
81	AAGAATAGAAACTGGTTACACCCCAATCGATATTCCTG	120
	C C T C T C C C	
121	TCGCTAACGCAATTCTTTGAGTGAATTGTTCCCGGTG	160
	CT G A G G C C C G C G A	
161	CTGGATTTGTGTTAGGACTAGTTGATATAATATGGGAAT	200
	G C T C C C C C T	
201	TTTGCGCCCTCTCAATGGGACGCATTCTTGTACAAATT	240
	C A T C G G G	
241	GAACAGTTAACCAACAAAGAATAGAAGAATTGCTAGGA	280
	G G C G G C G C	
281	ACCAAGCCATTCTAGATTAGAAGGACTAACGAAATCTTA	320
	G C G G T G C	
321	TCAAATTACGCAGAACATCTTTAGAGAGTGGGAAGCAGAT	360
	C C T GAGC C C	
361	CCTACTAACCCAGCATTAAAGAGAAGAGATGCGTATTCAAT	400
	C T C C C G A	
401	TCAATGACATGAACAGTGCCCTTACAACCGCTATTCTCT	440
	C C T G C A C A T	
441	TTTGCAGTTCAAAATTATCAAGTTCTCTTTATCAGTA	480
	G C C G C C C G G G	
481	TATGTTCAAGCTGCAAATTACATTATCAGTTTGAGAG	520
	C A T C T C C C A G C G C T C	
521	ATGTTTCAGTGGACAAAGGTGGGATTTGATGCCGC	560
	C A G C G C T	
561	GACTATCAATAGCTGTTATAATGATTAACCTAGGCTTATT	600
	A C C C C C C T G	
601	GGCAACTATACAGATTATGCTGTACGGCTGGTACAATACGG	640
	A C C C C C T T C T	
641	GATTAGAACGTGTATGGGGACCGGATTCTAGAGATTGGGT	680
	C G G C T T A	

FIG. 9A

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681	AAGGTATAATCAATTAGAAGAGAATTAAACACTAACTGTA T A C C G C G G C C A T	720
721	TTAGATATCGTGTCTGTTCCCGAATTATGATAGTAGAA G C T G T C C C C T C C	760
761	GATATCCAATTGAAACAGTTCCCAATTAAACAAGAGAAAT CC C T C T G C T C	800
801	TTATACAAACCCAGTATTAGAAAATTGATGGTAGTTT C T T C T G C C C C C	840
841	CGAGGCTCGGCTCAGGGCATAGAAAGAAGTATTAGGAGTC T T T C A T C G C T C C	880
881	CACATTTGATGGATATACTTAACAGTATAACCATCTATAC C C T G C T C	920
921	GGATGCTCATAGGGTTATTATTATTGGTCAGGGCATCAA C C A A G G C T A C G	960
961	ATAATGGCTTCTCCTGTAGGGTTTCGGGCCAGAATTCA C C A T A C A G C C G T	1000
1001	CTTTTCCGCTATATGGAACTATGGAAATGCAGCTCCACA C T C C C	1040
1041	ACAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATAGA C T C C	1080
1081	ACATTATCGTCCACTTATAGAAGACCTTTAATATAG C G T C G C C C	1120
1121	GGATAAATAATCAACAACTATCTGTTCTGACGGGACAGA T C C C G T C A	1160
1161	ATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTGTA G C C T T C T	1200
1201	TACAGAAAAAGCGGAACGGTAGATTGCTGGATGAAATAC G C T C T C C	1240
1241	CGCCACAGAATAACAACGTGCCACCTAGGCAAGGATTAG A C T C C T C	1280
1281	TCATCGATTAAGCCATGTTCAATGTTCGTTCAAGGCTT C C A G G C G C C C A C	1320
1321	AGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTTCT C C T C C G C C C	1360
1361	CTTGGATACATCGTAGTGCTGAATTAAATAATATAATTGC C G C C C C C	1400

FIG. 9B

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1401	ATCGGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC C	1440
1441	TTTCTTTTAATGGTTCTGTAATTCAGGACCAGGATTAC C C C C C	1480
1481	CTGGTGGGGACTTAGTTAGATTAAATAGTAGTGGAAATAA A C C C C C	1520
1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
1561	CCATCGACATCTACCAGATATCGAGTTCGTGTACGGTATG C A GA	1600
1601	CTTCTGTAACCCGATTCACCTAACGTTAATTGGGGTAA G T	1640
1641	TTCATCCATTTTCCAATACAGTACCACTACAGCTACAGCTACG C C T C	1680
1681	TCATTAGATAATCTACAATCAAGTGATTTGGTTATTTG C G C C C C	1720
1721	AAAGTGCCAATGCTTTACATCTCATTAGGTAAATATAAGT C C C C	1760
1761	AGGTGTTAGAAATTTAGTGGACTGCAGGAGTGATAATA G C T C	1800
1801	GACAGATTGAAATTATTCCAGTTACTGCAACACTCGAGG C G C	1840
1841	CTGAATATAATCTGGAAAGAGCGCAGAAGGCGGTGAATGC	1880
1881	GCTGTTACGTCTACAAACCAACTAGGGCTAAAACAAAT	1920
1921	GTAACGGATTATCATATTGATCAAGTGTCCAATTAGTTA	1960
1961	CGTATTTATCGGATGAATTGTCTGGATGAAAAGCGAGA	2000
2001	ATTGTCCGAGAAAGTCAAACATGCGAAGCGACTCAGTGAT	2040
2041	GAACGCAATTACTCCAAGATTCAAATTCAAAGACATTA	2080
2081	ATAGGCAACCAGAACGTGGGTGGGGCGGAAGTACAGGGAT	2120

FIG. 9C

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2121	TACCATCCAAGGAGGGGATGACGTATTTAAAGAAAATTAC	2160
2161	GTCACACTATCAGGTACCTTGATGAGTGCTATCCAACAT	2200
2201	ATTTGTATCAAAAAATCGATGAATCAAAATTAAAAGCCTT	2240
2241	TACCCGTTATCAATTAAGAGGGTATATCGAAGATAGTCAA	2280
2281	GACTTAGAAATCTATTTAATTCGCTACAATGCAAAACATG	2320
2321	AAACAGTAAATGTGCCAGGTACGGGTTCCCTATGGCCGCT	2360
2361	TTCAGCCAAAGTCCAATCGGAAAGTGTGGAGAGCCGAAT	2400
2401	CGATGCGCGCCACACCTTGAATGGAATCCTGACTTAGATT	2440
2441	GTTCGTGTAGGGATGGAGAAAAGTGTGCCCATTCGCA	2480
2481	TCATTTCTCCTTAGACATTGATGTAGGATGTACAGACTTA	2520
2521	AATGAGGACCTAGGTGTATGGTGATCTTAAGATTAAGA	2560
2561	CGCAAGATGGCACGCAAGACTAGGGATCTAGAGTTCT	2600
2601	CGAAGAGAAACCATTAGTAGGAGAAGCGCTAGCTCGTGT	2640
2641	AAAAGAGCGGAGAAAAATGGAGAGACAAACGTGAAAAT	2680
2681	TGGAATGGGAAACAAATATCGTTATAAGAGGGCAAAAGA	2720
2721	ATCTGTAGATGCTTATTTGTAACCTCTCAATATGATCAA	2760
2761	TTACAAGCGGATACGAATATTGCCATGATTGCGGCAG	2800
2801	ATAAACGTGTTCATAGCATTGAGAAGCTTATCTGCCTGA	2840

**FIG. 9D**

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2841	GCTGTCTGTGATTCCGGGTGTCAATGCGGCTATTTTGAA	2880
2881	GAATTAGAAGGGCGTATTTCACTGCATTCTCCCTATATG	2920
2921	ATGCGAGAAATGTCATTAAAAATGGTATTTAATAATGG	2960
2961	CTTATCCTGCTGGAACGTGAAAGGGCATGTAGATGTAGAA	3000
3001	GAACAAAACAACCAACGTTGGTCCTTGTGTTCCCGGAAT	3040
3041	GGGAAGCAGAAGTGTACAAAGAAGTTCGTGTCTGTCCGGG	3080
3081	TCGTGGCTATATCCTCGTGTACAGCGTACAAGGAGGGAA	3120
3121	TATGGAGAAGGTTGCGTAACCATTGAGATCGAGAACAA	3160
3161	ATACAGACGAACTGAAGTTAGCAACTGCGTAGAACAGAGGA	3200
3201	AATCTATCCAAATAAACACGGTAACGTGTAATGATTATACT	3240
3241	GTAAATCAAGAAGAATAACGGAGGTGCGTACACTCTCGTA	3280
3281	ATCGAGGATATAACGAAGCTCCTTCCGTACCGAGCTGATTA	3320
3321	TGCGTCAGTCTATGAAGAAAAATCGTATACAGATGGACGA	3360
3361	AGAGAGAATCCTTGTGAATTAAACAGAGGGTATAGGGATT	3400
3401	ACACGCCACTACCAGTTGGTTATGTGACAAAAGAATTAGA	3440
3441	ATACTTCCCAGAAACCGATAAGGTATGGATTGAGATTGGA	3480
3481	GAAACGGAAGGAACATTTATCGTGGACAGCGTGGATTAC	3520
3521	TCCTTATGGAGGAA 3534	

FIG. 9E

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1	ATGGATAACAATCCGAACATCAATGAATGCATTCTTATA	40
	C C A C A C	
41	ATTGTTAACGTAACCCCTGAAGTAGAAGTATTAGGTGGAGA	80
	C C G A T C T	
81	AAGAATAGAAACTGGTTACACCCCAATCGATATTCCTTG	120
	C C T C T C C C	
121	TCGCTAACGCAATTCTTTGAGTGAATTGTTCCCGGTG	160
	CT G A G G C C C G C G A	
161	CTGGATTTGTGTTAGGACTAGTTGATATAATATGGGAAT	200
	G C T C C C C C T	
201	TTTGGTCCCTCTCAATGGGACGCATTCTGTACAAATT	240
	C A T C G G G	
241	GAACAGTTAACCAACAAAGAATAGAAGAATTGCTAGGA	280
	G G C G G C G C	
281	ACCAAGCCATTCTAGATTAGAAGGACTAAGCAATCTTA	320
	G C G G T G C	
321	TCAAATTACGCAGAATCTTTAGAGAGTGGGAAGCAGAT	360
	C C T GAGC C C	
361	CCTACTAACCGCATTAAGAGAAGAGATGCGTATTCAAT	400
	C T C C C G A	
401	TCAATGACATGAACAGTGCCCTACAAACCGCTATTCTCT	440
	C C T G C A C A T	
441	TTTGCAGTTCAAAATTATCAAGTTCTCTTATCAGTA	480
	G C C G C C C G G	
481	TATGTTCAAGCTGCAAATTACATTTATCAGTTGAGAG	520
	C A T C T C C C A G C G T C	
521	ATGTTTCAGTGTGGACAAAGGTGGGATTTGATGCCGC	560
	C A G C G C T	
561	GACTATCAATAGTCGTTATAATGATTAACTAGGCTTATT	600
	A C C C C C T G	
601	GGCAACTATACAGATTATGCTGTACGCTGGTACAATACGG	640
	A C C C C T T C T	

FIG. 10A

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641	GATTAGAACGTGTATGGGGACCGGATTCTAGAGATTGGGT C G G C T T A	680
681	AAGGTATAATCAATTAGAAGAGAATTAACACTAACTGTA T A C C G C G G C C A T	720
721	TTAGATATCGTTGCTCTGTTCCCGAATTATGATAGTAGAA G C T G T C C C C T C C	760
761	GATATCCAATTGAAACAGTTCCCAATTAACAAGAGAAAT CC C T C T G C T C C	800
801	TTATACAAACCCAGTATTAGAAAATTTGATGGTAGTTT C T T C T G C C C C C	840
841	CGAGGCTCGGCTCAGGGCATAGAAAGAAGTATTAGGAGTC T T T C A T C G C T C C C	880
881	CACATTTGATGGATATACTTAACAGTATAACCATCTATAC C C T G C T C C	920
921	GGATGCTCATAGGGTTATTATTATTGGTCAGGGCATCAA C C A A G G C T A C G	960
961	ATAATGGCTTCTCCTGTAGGGTTTGGGGCCAGAATTCA C C A T A C A G G C G T	1000
1001	CTTTCCGCTATATGGAACTATGGAAATGCAGCTCCACA C T C C C	1040
1041	ACAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATAGA C T C C	1080
1081	ACATTATCGTCCACTTATATAGAACGACCTTTAATATAG C G T C G C C C C	1120
1121	GGATAAATAATCAACAACTATCTGTTCTGACGGGACAGA T C C C G T C A	1160
1161	ATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTGTA G C C T T C T	1200
1201	TACAGAAAAAGCGGAACGGTAGATTGCTGGATGAAATAC G C T C T C C	1240
1241	CGCCACAGAATAACAACGTGCCACCTAGGCAAGGATTTAG A C T C C T C	1280
1281	TCATCGATTAAGCCATGTTCAATGTTGTTCAAGGCTTT C C A G G C G C C C A C	1320
1321	AGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTTCT C C T C C G C C C	1360

FIG. 10B

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1361	CTTGGATACATCGTAGTGCTGAATTAAATAATATAATTGC C G C C C C C	1400
1401	ATCGGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC C	1440
1441	TTTCTTTAATGGTTCTGTAATTTCAGGACCAGGATTTC C C C C C C	1480
1481	CTGGTGGGGACTTAGTTAGATTAAATAGTAGTGGAAATAA A C C C C C	1520
1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
1561	CCATCGACATCTACCAGATATCGAGTTCGTGTACGGTATG C A G A	1600
1601	CTTCTGTAACCCCGATTCACCTAACGTTAATTGGGGTAA G T	1640
1641	TTCATCCATTTCACAGTACAGCTACAGCTACAGCTACG C C T C	1680
1681	TCATTAGATAATCTACAATCAAGTGATTTGGTTATTTG C G C C C C	1720
1721	AAAGTGCCAATGCTTTACATCTTCATTAGGTAATATAGT C C C C	1760
1761	AGGTGTTAGAAATTAGTGGACTGCAGGAGTGATAATA G C T C	1800
1801	GACAGATTGAATTATTCCAGTTACTGCAACACTCGAGG C G C	1840
1841	CTGAATATAATCTGGAAAGAGCGCAGAAGGCAGGTGAATGC	1880
1881	GCTGTTACGTCTACAAACCAACTAGGGCTAAAAACAAAT G C C C G C	1920
1921	GTAACGGATTATCATATTGATCAAGTGTCCAATTAGTTA G C G G	1960
1961	CGTATTTATCGGATGAATTGTCTGGATGAAAGCGAGA C CC CAGC G C	2000
2001	ATTGTCCGAGAAAGTCAAACATGCGAAGCGACTCAGTGAT	2040
2041	GAACGCAATTACTCCAAGATTCAAATTCAAAGACATTA	2080

FIG.10C

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2081	ATAGGCAACCAGAACGTGGGTGGGCAGGAAGTACAGGGAT	2120
2121	TACCATCCAAGGAGGGATGACGTATTTAAAGAAAATTAC G T C G C G G C	2160
2161	GTCACACTATCAGGTACCTTGATGAGTGCTATCCAACAT	2200
2201	ATTTGTATCAAAAAATCGATGAATCAAAATTAAAAGCCTT CC C C G G C G C G G	2240
2241	TACCCGTTATCAATTAAGAGGGTATATCGAAGATAAGTCAA	2280
2281	GACTTAGAAATCTATTTAATTGCTACAATGCAAAACATG C C G C C C	2320
2321	AAACAGTAAATGTGCCAGGTACGGGTTCCCTATGGCCGCT	2360
2361	TTCAGCCAAAGTCCAATCGGAAAGTGTGGAGAGCCGAAT	2400
2401	CGATGCGGCCACACCTTGAATGGAATCCTGACTTAGATT	2440
2441	GTTCGTGTAGGGATGGAGAAAAGTGTGCCCATTCGCA	2480
2481	TCATTTCTCCTTAGACATTGATGTAGGATGTACAGACTTA	2520
2521	AATGAGGACCTAGGTGTATGGGTGATCTTAAGATTAAGA	2560
2561	CGCAAGATGGGCACGCAAGACTAGGGATCTAGAGTTCT	2600
2601	CGAAGAGAAACCATTAGTAGGAGAAGCGCTAGCTCGTGT	2640
2641	AAAAGAGCGGAGAAAAATGGAGAGACAAACGTGAAAAT G G	2680
2681	TGGAATGGAAACAAATATCGTTATAAAGAGGCAAAAGA G C C C C	2720
2721	ATCTGTAGATGCTTATTTGAACTCTCAATATGATCAA	2760
2761	TTACAAGCGGATACGAATATTGCCATGATTGCGGCAG	2800

FIG. 10D

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2801	ATAAACGTGTTCATAGCATTGAGAAGCTTATCTGCCTGA	2840
2841	GCTGTCTGTGATTCCGGGTGTCAATGCGGCTATTTTGAA	2880
2881	GAATTAGAAGGGCGTATTTCACTGCATTCTCCCTATATG C C	2920
2921	ATGCGAGAAATGTCATTAAAATGGTGATTTAATAATGG C C C G C C C C	2960
2961	CTTATCCTGCTGGAACGTGAAAGGGCATGTAGATGTAGAA	3000
3001	GAACAAAACAACCAACGTTGGTCCTTGTGTTCCGGAAT	3040
3041	GGGAAGCAGAAGTGTCAACAAGAAGTTCGTGTCTGTCGGG	3080
3081	TCGTGGCTATATCCTTCGTGTACAGCGTACAAGGAGGG	3120
3121	TATGGAGAAGGTTGCGTAACCATTGAGATCGAGAACAA	3160
3161	ATACAGACGAACTGAAGTTAGCAACTGCGTAGAAGAGGA	3200
3201	AATCTATCCAAATAACACGGTAACGTGTAATGATTATACT	3240
3241	GTAAATCAAGAAGAATACGGAGGTGCGTACACTTCTCGTA	3280
3281	ATCGAGGATATAACGAAGCTCCTCCGTACCGAGCTGATTA	3320
3321	TGCGTCAGTCTATGAAGAAAAATCGTATACAGATGGACGA	3360
3361	AGAGAGAATCCTGTGAATTAAACAGAGGGTATAGGGATT	3400
3401	ACACGCCACTACCAGTTGGTTATGTGACAAAAGAATTAGA	3440
3441	ATACTTCCCAGAAACCGATAAGGTATGGATTGAGATTGGA	3480
3481	GAAACGGAAGGAACATTTATCGTGGACAGCGTGGATTAC	3520
3521	TCCTTATGGAGGAA 3534	

FIG. 10E

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1	ATGGATAACAATCCGAACATCAATGAATGCATTCTTATA	40
	C C A C A C	
41	ATTGTTAACGTAACCTGAAGTAGAAGTATTAGGTGGAGA	80
	C C G A T C T	
81	AAGAATAGAAACTGGTTACACCCCAATCGATATTCCTTG	120
	C C T C T C C C	
121	TCGCTAACGCAATTCTTTGAGTGAATTGTTCCGGTG	160
	CT G A G G C C C G C G A	
161	CTGGATTGTGTTAGGACTAGTTGATATAATATGGGAAT	200
	G C T C C C C C T	
201	TTTGGTCCCTCTCAATGGGACGCATTCTTGACAAATT	240
	C A T C G G G	
241	GAACAGTTAACCAACAAAGAATAGAAGAATTGCTAGGA	280
	G G C G G C G C	
281	ACCAAGCCATTCTAGATTAGAAGGACTAACGCAATCTTA	320
	G C G G T G C	
321	TCAAATTACGCAGAACATCTTTAGAGAGTGGGAAGCAGAT	360
	C C T GAGC C C	
361	CCTACTAACCCAGCATTAAGAGAAGAGATGCGTATTCAAT	400
	C T C C C G A	
401	TCAATGACATGAACAGTGCCTTACAACCGCTATTCT	440
	C C T G C A C AT	
441	TTTGCAGTTCAAAATTATCAAGTTCCCTTTTATCAGTA	480
	G C C G C C C G G C G C G	
481	TATGTTCAAGCTGCAAATTACATTATCAGTTGAGAG	520
	C A T C T C C C A G C G C T C	
521	ATGTTTCAGTGTGTTGGACAAAGGTGGGATTGATGCCGC	560
	C AGC G C T	
561	GACTATCAATAGTCGTTATAATGATTAACTAGGCTTATT	600
	A C C C C C C T G	
601	GGCAACTATACAGATTATGCTGTACGCTGGTACAATACGG	640
	A C C C C C T T C T	
641	GATTAGAACGTGTATGGGGACCGGATTCTAGAGATTGGGT	680
	C G G C T T A	

FIG. 11A

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681	AAGGTATAATCAATTAGAAGAGAATTAAACACTAACTGTA T A C C G C G G C C A T	720
721	TTAGATATCGTGTCTGTTCCCGAATTATGATAGTAGAA G C T G T C C C C C T C C	760
761	GATATCCAATTGAAACAGTTCCAATTAAACAAGAGAAAT CC C T C T G C T C	800
801	TTATACAAACCCAGTATTAGAAAATTGATGGTAGTTT C T T C T G C C C C C C	840
841	CGAGGCTCGGCTCAGGGCATAGAAAGAAGTATTAGGAGTC T T T C A T C G C T C C	880
881	CACATTTGATGGATATACTTAACAGTATAACCATCTATAC C C T G C T C	920
921	GGATGCTCATAGGGTTATTATTATTGGTCAGGGCATCAA C C A A G G C T A C G	960
961	ATAATGGCTTCTCTGTAGGGTTTCGGGCCAGAATTCA C C A T A C A G C C G T	1000
1001	CTTTCCGCTATATGGAACTATGGAAATGCAGCTCCACA C T C C C	1040
1041	ACAACGTATTGTTGCTCAACTAGGTCAAGGGCGTGTATAGA C T C C	1080
1081	ACATTATCGTCCACTTATATAGAAGACCTTTAATATAG C G T C G C C C	1120
1121	GGATAAATAATCAACAACTATCTGTTCTGACGGGACAGA T C C C G T C A	1160
1161	ATTTGCTTATGGAACCTCCTCAAATTGCCATCCGCTGTA G C C T T C T	1200
1201	TACAGAAAAGCGGAACGGTAGATTCGCTGGATGAAATAC G C T C T C C	1240
1241	CGCCACAGAATAACAACGTGCCACCTAGGCAAGGGATTTAG A C T C C T C	1280
1281	TCATCGATTAAGCCATGTTCAATGTTCGTTCAAGGCTT C C A G G C G C C C A C	1320
1321	AGTAATAGTAGTGTAAAGTATAATAAGAGCTCCTATGTTCT C C T C C G C C C	1360
1361	CTTGGATACATCGTAGTGCTGAATTAAATAATTGC C G C C C C C	1400

FIG. 11B

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1401	ATCGGATAGTATTACTCAAATCCCTGCAGTGAAGGGAAAC	1440
	C	
1441	TTTCTTTTAATGGTTCTGTAATTCAGGACCAGGATTAC	1480
	C C C C	C
1481	CTGGTGGGGACTTAGTTAGATTAAATAGTAGTGGAAATAA	1520
	A C C C C C	
1521	CATTCAGAATAGAGGGTATATTGAAGTTCCAATTCACTTC	1560
1561	CCATCGACATCTACCAGATATCGAGTTCGTGTACGGTATG	1600
	C A GA	
1601	CTTCTGTAACCCCGATTCACCTCAACGTTAATTGGGGTAA	1640
	G T	
1641	TTCATCCATTTCACAGTACAGTACAGCTACAGCTACG	1680
	C C T C	
1681	TCATTAGATAATCTACAATCAAGTGATTTGGTTATTTG	1720
	C G C C C C	
1721	AAAGTCCAATGCTTTACATCTCATTAGGTAAATATAGT	1760
	CC C C C	
1761	AGGTGTTAGAAATTTAGTGGGACTGCAGGAGTGATAATA	1800
	G C T C	
1801	GACAGATTGAAATTATTCCAGTTACTGCAACACTCGAGG	1840
	C G C	
1841	CTGAATATAATCTGGAAAGAGCGCAGAAGGCAGTGAATGC	1880
	G C C T G C T C	
1881	GCTGTTACGTCTACAAACCAACTAGGGCTAAAAACAAAT	1920
	C C C C T G T C T G T C	
1921	GTAACGGATTATCATATTGATCAAGTGTCCAATTAGTTA	1960
	T T C C C C C G C	
1961	CGTATTTATCGGATGAATTGTCTGGATGAAAAGCGAGA	2000
	C CC TAGC G C C C C G T	
2001	ATTGTCCGAGAAAGTCAAACATGCGAAGCGACTCAGTGAT	2040
	C C T C C T C C	
2041	GAACGCAATTACTCCAAGATTCAAATTCAAAGACATTA	2080
	GA G C CT G C C C C C	
2081	ATAGGCAACCAGAACGTGGGTGGGGCGGAAGTACAGGGAT	2120
	C G T T C C	

FIG. 11C

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2121	TACCATCCAAGGAGGGATGACGTATTTAAAGAAAATTAC C C C T G C G G C	2160
2161	GTCACACTATCAGGTACCTTGATGAGTGCTATCCAACAT C C C A T C C C T C	2200
2201	ATTTGTATCAAAAAATCGATGAATCAAAATTAAAAGCCTT C C G G G C C C	2240
2241	TACCCGTTATCAATTAAGAGGGTATATCGAAGATAGTCAA C A G C T C C C C	2280
2281	GACTTAGAAATCTATTTAATCGCTACAATGCAAAACATG C T C C G C A G C G C	2320
2321	AAACAGTAAATGTGCCAGGTACGGGTTCCCTATGGCCGCT G C G C T C C A	2360
2361	TTCAGCCAAAGTCCAATCGAAAGTGTGGAGAGCCGAAT T T C T G T C	2400
2401	CGATGCGGCCACACCTTGAATGGAATCCTGACTTAGATT A T G G C	2440
2441	GTTCGTGTAGGGATGGAGAAAAGTGTGCCCATTCGCA C C C C G C T	2480
2481	TCATTTCTCCTTAGACATTGATGTAGGATGTACAGACTTA C G C G T C G	2520
2521	AATGAGGACCTAGGTGTATGGGTGATCTTAAGATTAAGA C A C C C C	2560
2561	CGCAAGATGGGCACGCAAGACTAGGGAATCTAGAGTTCT C C A T C C C T	2600
2601	CGAAGAGAAACCATTAGTAGGAGAAGCGCTAGCTCGTGTG G C T T C	2640
2641	AAAAGAGCGGAGAAAAATGGAGAGACAAACGTAAAAAT G A G G G G G C	2680
2681	TGGAATGGAAACAAATATCGTTATAAGAGGGCAAAAGA C T C C G C	2720
2721	ATCTGTAGATGCTTATTTGTAAACTCTCAATATGATCAA G C G G C G C G	2760
2761	TTACAAGCGGATACGAATATTGCCATGATTCATGCGGCAG G C C C C C C C	2800
2801	ATAAACGTGTTCATAGCATTGAGAAGCTTATCTGCCTGA C G C T G C T	2840

FIG. 11D

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2841	GCTGTCTGTGATTCCGGGTGTCAATGCGGCTATTTTGAA T C C T G C T C C C G	2880
2881	GAATTAGAAGGGCGTATTTCACTGCATTCTCCCTATATG C T G A C T C T G C	2920
2921	ATGCGAGAAAATGTCATTAATGGTGATTTAATAATGG C C C G C C C C	2960
2961	CTTATCCTGCTGAAACGTGAAAGGGCATGTAGATGTAGAA C CAG T T G C G G G	3000
3001	GAACAAAACAACCAACGTTCGGTCTTGTGTTCCGGAAT G T G C G G T G	3040
3041	GGGAAGCAGAAGTGTACAAGAAGTTCGTGTCTGCCGG T C G A A A	3080
3081	TCGTGGCTATATCCTCGTGTACAGCGTACAAGGAGGG A A C T C G C T	3120
3121	TATGGAGAAGGTTGCGTAACCATTGAGATCGAGAAC C T G G C C	3160
3161	ATACAGACGAACTGAAGTTAGCAACTGCGTAGAAGAGGA C C G T CTC C G A	3200
3201	AATCTATCCAAATAACACGGTAACGTGTAATGATTATACT C C C T T C C C C	3240
3241	GTAAATCAAGAAGAATACGGAGGTGCGTACACTCTCGTA G G G C C AGC	3280
3281	ATCGAGGATATAACGAAGCTCCTCCGTACCGCTGATTA CA T C T T C	3320
3321	TGCGTCAGTCTATGAAGAAAATCGTATACAGATGGACGA C C G C G G C C C A	3360
3361	AGAGAGAATCCTTGTGAATTAAACAGAGGGTATAGGGATT C T C C G C T C C	3400
3401	ACACGCCACTACCAGTTGGTTATGTGACAAAAGAATTAGA A T C T C G G C T	3440
3441	ATACTTCCCAGAAACCGATAAGGTATGGATTGAGATTGGA G T T G C A G C C T	3480
3481	GAAACGGAAAGGAACATTTATCGTGGACAGCGTGGATTAC C G C C G C T	3520
3521	TCCTTATGGAGGAA 3534 T G	

FIG. 11E

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1	ATGACTGCAGATAATAATACGGAAGCACTAGATAGCTCTA C C C C C C C C T	40
41	CAACAAAAGATGTCATTCAAAAAGGCATTCGGTAGTAGG C T G T C G G T C T G	80
81	TGATCTCCTAGGCGTAGTAGGTTCCCGTTGGTAGCG A C T G G T A T C C C	120
121	CTTGTTCGTTTATACAAACTTTAAATACTATTTGGC C GAGC C C C C C	160
161	CAAGTGAAGACCCGTGGAAGGCCTTATGGAACAAAGTAGA C G T A A C G T	200
201	AGCATTGATGGATCAGAAATAGCTGATTATGCAAAAAT TC T G T A C G C	240
241	AAAGCTCTTGCAGAGTTACAGGGCCTCAAAATAATGTCG G T G A C C G C G	280
281	AAGATTATGTGAGTCATTGAGTCATGGCAAAAAATCC G C C T C C A G G G C	320
321	TGTGAGTTCACGAAATCCACATAGCCAGGGCGGATAAGA T C C A T C A T A C	360
361	GAGCTGTTTCTCAAGCAGAAAGTCATTTCGTAATTCAA T C C T C C C A A C	400
401	TGCCTTCGTTGCAATTCTGGATACGAGGTTCTATTCT AGC T C C T T C	440
441	AACAAACATATGCACAAGCTGCCAACACACATTATTTA C T C T C C G C C	480
481	CTAAAAGACGCTCAAATTATGGAGAAGAATGGGATACG T G C G	520
521	AAAAAGAAGATATTGCTGAATTATAAAAGACAATCAA G G C G C C G C T T	560
561	ACTTACGCAAGAATATACTGACCATTGTGTCAAATGGTAT G C C G C G	600
601	AATGTTGGATTAGATAAAATTAAGAGGTTCATCTTATGAAT C T C C G C C T C C G	640
641	CTTGGGTAAACTTAACCGTTATCGCAGAGAGATGACATT G C A A C A G C	680

FIG. 12A

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681	AACAGTATTAGATTAATTGCACTATTCATTGTATGAT	720
	G T G C C C T C C C C	
721	GTTCGGCTATAACCCAAAAGAAGTTAAAACCGAATTAACAA	760
	GA A C G G T G C T C	
761	GAGACGTTTAACAGATCCAATTGTCGGAGTCAACAAACCT	800
	GC C T C T	
801	TAGGGGCTATGGAACAAACCTTCTCTAAATATAGAAAATTAT	840
	T T AGC C C C	
841	ATTCGAAAACCACATCTATTGACTATCTGCATAGAATTCA	880
	A G C C T C	
881	AATTCACACGGTTCCAACCAGGATATTATGGAAATGAC	920
	C AA T C T C	
921	CTCTTCATTATTGGTCCGGTAATTATGTTCAACTAGA	960
	C C C C C	
961	CCAAGCATAGGATCAAATGATATAATCACATCTCCATTCT	1000
	T T C C C	
1001	ATGGAAATAATCCAGTGAACCTGTACAAAATTAGAATT	1040
	T C G G G C C T G	
1041	TAATGGAGAAAAAGTCTATAGAGCCGTAGCAAATACAAAT	1080
	C C C G C C C	
1081	CTTGCGGTCTGGCCGTCCGCTGTATATTAGGTGTTACAA	1120
	C T G A A T C C C	
1121	AAGTGGAAATTAGCCAATATAATGATCAAACAGATGAAGC	1160
	G G T G C G C G	
1161	AAGTACACAAACGTACGACTCAAAAGAAATGTTGGCGCG	1200
	C C C G T C T C A	
1201	GTCAGCTGGATTCTATCGATCAATTGCCCTCAGAAACAA	1240
	TCT C C	
1241	CAGATGAACCTCTAGAAAAGGGATATAGCCATCAACTCAA	1280
	C AT G G C C C T	
1281	TTATGTAATGTGCTTTTAATGCAGGGTAGTAGAGGAACA	1320
	C G C G A T C C G C	
1321	ATCCCAGTGTAACTTGGACACATAAAAGTGTAGACTTT	1360
	T G C C G T C G C	
1361	TTAACATGATTGATTGAAAAAAATTACACAACTTCCGTT	1400
	C C AGC G G C T C	

FIG.12B

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1401	AGTAAAGGCATATAAGTTACAATCTGGTGCTTCGTTGTC	1440
	G G A C C C G	
1441	GCAGGTCCCTAGGTTACAGGAGGAGATATCATTCAATGCA	1480
	C A C T T C C G	
1481	CAGAAAATGGAAGTGC GG CAACTATTTACGTTACACCGGA	1520
	G C C C A T C G T	
1521	TGTGTCGTACTCTCAAAAATATCGAGCTAGAATTCAATTAT	1560
	T G G C A G A C T C	
1561	GCTTCTACATCTCAGATAACATTTACACTCAGTTAGACG	1600
	A CAGC C C C C G T	
1601	GGGCACCATTAAATCAATACTATTCGATAAAACGATAAA	1640
	A C C C G T C T C G C C	
1641	TAAAGGAGACACATTAACGTATAATTCAATTAAATTAGCA	1680
	C T T C C A C AGC C C G	
1681	AGTTTCAGCACACCATTCAATTATCAGGGAAATAACTTAC	1720
	T C C C C T C T	
1721	AAATAGGCGTCACAGGATTAAGTGCCTGGAGATAAGTTA	1760
	G C C T C C C C C	
1761	TATAGACAAAATTGAATTATTCCAGTGAAT	1791
	C C G G C C C	

FIG.12C

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1	ATG	AATAATGTATTGAATAGTGGAAAGAACAACTATTT	40
	GAC	C C C C CTC T C C	
41	GTGATGCGTATAATGTAGTAGCCCATGATCCATTAGTT	80	
	C C A C C C G T C C C		
81	TGAACATAAAATCATTAGATACCATCCAAAAGAACATGGATG	120	
	C C GAGCC C C T T G G G		
121	GAGTGGAAAAGAACAGATCATAGTTATATGTAGCTCCTG	160	
	A C T T C CTC C C C C A		
161	TAGTCGGAACTGTGTCTAGTTTTGCTAAAGAAAAGTGGG	200	
	G T A C C C C T C G C		
201	GAGTCTTATTGGAAAAAGGATATTGAGTGAATTATGGGGG	240	
	CTC C C C T C TCC C C T		
241	ATAATATTCCTAGTGGTAGTACAAATCTAATGCAAGATA	280	
	C C ATC GTCC T C C		
281	TTTTAAGGGAGACAGAACAAATTCTAAATCAAAGACTTAA	320	
	C G C G T C C C G C T C		
321	TACAGATACCCCTTGCTCGTGTAAATGCAGAATTGATAGGG	360	
	C T T G A A C C T G C T		
361	CTCCAAGCGAATATAAGGGAGTTAACAAACAAAGTAGATA	400	
	A C T C T C C G G C		
401	ATTTTTAAACCTACTCAAAACCTGTTCTTATCAAT	440	
	C C G T A G T G C T C		
441	AACTTCTCGGTTAACATACAATGCAGCAATTATTCCTAAAT	480	
	C C G C T C C C C C		
481	AGATTACCCCAGTTCCAGATACAAAGGATACCAGTTGTTAT	520	
	G T T T C C		
521	TATTACCTTATTGCACAGGCAGCCAATATGCATCTTC	560	
	T C T A C C T T C C T G		
561	TTTTATTAGAGATGTTATTCTTAATGCAGATGAATGGGGT	600	
	C C A C T C G C C C T C A		
601	ATTTCAGCAGCAACATTACGTACGTACGATTACCTGA	640	
	C T C T C T A G A C A C T		
641	GAAATTATAACAGAGATTATTCTAATTATTGTATAAATAC	680	
	G C C T C T C C C C C C		

FIG. 13A

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681	GTATCAAAC TGC GTTAGAGGGTTAACACCCGTTACAC	720
	T G C C T A C C T T A G C T	
721	GATATGTTAGAATTAGAACATATATGTTTAAATGTAT	760
	C C T G C G C C C C T C G	
761	TTGAATATGTATCCATTGGTCATTGTTAAATATCAGAG	800
	G C C A G A G T C C C G C	
801	TCTTATGGTATCTTCTGGCGCTAATTATATGCTAGCGGT	840
	C T G G C A C C C C C T C T C	
841	AGTGGACCACAGCAGACACAATCATTACAGCACAAAAT	880
	A T G A G C C T G	
881	GGCCATT T T T A T T C T C T T C C A G T T A A T C G A A T T A	920
	C G A G C T G C C C C C	
921	TATATTATCTGGTATTAGTGGTACTAGGCTTCTATTACC	960
	C T C C A G C T C G C A C C C A	
961	TTCCCTAATATTGGTGGTTACCGGGTAGTACTACAACTC	1000
	T C C A C T A C T C C C	
1001	ATT CATTGAATAGTGCCAGGGTTAATTATAGCGGAGGAGT	1040
	A G C C T T	
1041	TTCATCTGGTCTCATAGGGCGACTAATCTCAATCACAAC	1080
	C A G C A T G T T A C T G G C	
1081	TTTAATTGCAGCACGGTCCTCCCTCCTTATCAACACCAT	1120
	C T C T G A C G A G C G	
1121	TTGTTAGAAGTTGGCTGGATTCAAGGTACAGATCGAGAGGG	1160
	G G T C C T C A G C T C A	
1161	CGTTGCTACCTCTACGAATTGGCAGACAGAACGAACTCTTCAA	1200
	A A C A C G G C	
1201	ACAACTTTAAGTTAAGGTGTGGTCTTTTCAGCCGTG	1240
	C C T C C T A C T A	
1241	GAAATTCAAAC T A T T C C C A G A T T A T T T A T C C G T A A T A T	1280
	G C T C C C T A G C	
1281	TTCTGGGTTCTTCTAGTTATTAGAAACGAAGATCTAACAC	1320
	C T C C C C G T C C C C	
1321	AGACCGTTACACTATAACCAAATAAGAAATATAGAAAGTC	1360
	C T A C T T C G T G C C C G T C	
1361	CTTCGGGAACACCTGGTGGAGCACGGGCCTATTGGTATC	1400
	A C T T A A T A A T C C C G	

FIG.13B

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1401	TGTGCATAACAGAAAAATAATATCTATGCCGCTAATGAA	1440
	C G G C C C T C C C G	
1441	AATGGTACTATGATCCATTGGCGCCAGAAGATTATACAG	1480
	C C T C C T A C T	
1481	GATTTACTATATGCCAATACATGCCACTCAAGTGAATAA	1520
	C C C T C T C C	
1521	TCAAACCTCGAACATTTATTCTGAAAAATTGGAAATCAA	1560
	G A C C C C C G C	
1561	GGTGATTCTTAAGATTGAAACAAAGCAACACGACAGCTC	1600
	C G G C G T C T C A	
1601	GTTATACGCTTAGAGGGAATGGAAATAGTTACAATCTTA	1640
	G C T T G C C C C	
1641	TTTAAGAGTATCTCAATAGGAAATTCAACTATTGAGTT	1680
	C G T A G C C T C C C C T	
1681	ACTATAAACGGTAGAGTTACTGTTCAAATGTTAATA	1720
	C C A C T G C	
1721	CCACTACAAATAACGATGGAGTTAATGATAATGGAGCTCG	1760
	T A G C T C C C C A	
1761	TTTTCAGATATTAATATCGGTAATATAGTAGCAAGTGAT	1800
	A C A G C C C T C C C G C T C C	
1801	AATACTAATGTAACGCTAGATATAATGTGACATTAAACT	1840
	C C T T T G C C C C T	
1841	CCGGTACTCCATTGATCTCATGAATATTATGTTGTGCC	1880
	T A C C C	
1881	AACTAATCTTCCACCACTTAT	1902
	C C T T G C	

FIG. 13C

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1	ATGGAGGAAAATAATCAAAATCAATGCATACCTTACAATT	40
	G C C C T A C	
41	GTTTAAGTAATCCTGAAGAAGTACTTTGGATGGAGAACG	80
	C G C A G T G C T	
81	GATATCAACTGGTAATTCAATTGATATTCCTCTGTCA	120
	C T C T C C C C C T C	
121	CTTGTTCAGTTCTGGTATCTAACTTTGTACCAGGGGAG	160
	T G C C A G C C G T T	
161	GATTTTAGTTGGATTAATAGATTTGTATGGGAATAGT	200
	G C C T C C T C C C T C	
201	TGGCCCTTCTCAATGGATGCATTTCTAGTACAAATTGAA	240
	T A C G G G	
241	CAATTAATTAATGAAAGAATAGCTGAATTTGCTAGGAATG	280
	G G C C G G C G C C C C	
281	CTGCTATTGCTAATTTAGAAGGATTAGGAAACAATTCAA	320
	C C C G G C T C	
321	TATATATGTGGAAAGCATTAAAGAATGGGAAGAAGATCCT	360
	C C G C C G G C G C	
361	AATAATCCAGAAACCAGGACCAGAGTAATTGATCGCTTC	400
	C G C C T G G G C C A A C A	
401	GTATACTTGATGGGCTACTTGAAAGGGACATTCCTTCGTT	440
	A C T G C C C T G G A T C A C	
441	TCGAATTCTGGATTGAAAGTACCCCTTTATCCGTTTAT	480
	C A C C C T T C G G C	
481	GCTCAAGCGGCCAATCTGCATCTAGCTATATTAGAGATT	520
	A T T C C C C C T C A	
521	CTGTAATTTGGAGAAAGATGGGATTGACAACGATAAA	560
	G C C G G C T C	
561	TGTCAATGAAAATATAATAGACTAATTAGGCATATTGAT	600
	C G T C C T C C C C	
601	GAATATGCTGATCACTGTGCAAATACGTATAATGGGGAT	640
	G C C C T C C C C T C	
641	TAAATAATTACCGAAATCTACGTATCAAGATTGGATAAC	680
	G C C C T G T T	
681	ATATAATCGATTACGGAGAGACTTAACATTGACTGTATTA	720
	C C C A G G A G C C C A T G	

FIG. 14A

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721	GATATGCCGCTTCTTC C T A C G C	760
761	ATCCAATTCA G C A G T C A C	800
801	TACGGACCC T C T C C C T G A A G	840
841	GTAGCTCAATTACCTACT C C C T C A C C T C	880
881	GAATTAGAAATCCTCATT T C G C A C G C C C C	920
921	TACAATCTTACGGATTGGTT T C C C C G T C C	960
961	TATTGGGGAGGACATCGAG T C A G C C C T C T T	1000
1001	GTGGTAACATAACATCTCCT G T C C C T A	1040
1041	CCAGGAGCCTCCAAGATCCTT A C T A G T C C C C T A C	1080
1081	TTTAGGACTTATCAAATCCTACT C A C G T C C G A G C C	1120
1121	AACCTTGGCCAGCGCCACC T T C C C T A	1160
1161	AGGAGTAGAATTCTACACCT G C T G C T C C T C	1200
1201	CGAGGAAGAGGTACGGTTGATT A T A C C G C C C A	1240
1241	CTGAGGATAATAGTGTGCCAC A C C C A G C C C T C C	1280
1281	TCGTTTATGTCATGCAACT C A G G C C C C G G C T C T	1320
1321	CCTTTTTAACAACTGGTGTAGT A C C C T A A T G C A T	1360
1361	GTAGTGCAACTCTTACAAATACA T C T C C G	1400

FIG. 14B

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1401	TAATCAAATACCTTAGTCAAAGGATTTAGAGTTGGGG C C A G C G T C C T G A	1440
1441	GGCACCTCTGTCATTACAGGACCAGGATTTACAGGAGGG A T C C C T	1480
1481	ATATCCTCGAAGAAATACCTTGTTGATTTGTATCTCT T A C T C C G A G C	1520
1521	ACAAGTCAATATTAATTACCAATTACCCAAAGATAACCGT C T C C C T T T	1560
1561	TTAAGATTCGTTACGCTTCCAGTAGGGATGCACGAGTTA C C G A TTCCC T C T A C	1600
1601	TAGTATTAACAGGAGCGGCATCCACAGGAGTGGGAGGCCA C G C C C A T T C T C T C T A	1640
1641	AGTTAGTGTAAATATGCCTCTTCAGAAAATATGGAAATA CTCC G C A C G G C	1680
1681	GGGGAGAACTTAACATCTAGAACATTAGATATACCGATT C G C G C C C C	1720
1721	TTAGTAATCCTTTTCATTAGAGCTAATCCAGATATAAT CTC C CAGT CC T C C T C C	1760
1761	TGGGATAAGTGAACAAACCTCTATTGGTGCAGGTTCTATT C T C C A T AGC C	1800
1801	AGTAGCGGTGAACCTTATATAGATAAAATTGAAATTATTC TCATCT C T G C T C G G C	1840
1841	TAGCAGATGCAACATTTGAAGCAGAATCTGATTTAGAAAG T C C T C C G T G A C A C C T G	1880
1881	AGCACAAAAGCGGTGAATGCCCTGTTACTTCTTCCAAT C G T C C C C C C A	1920
1921	CAAATCGGGTTAAAACCGATGTGACGGATTATCATATTG G C T C G T A C T T C C	1960
1961	ATCAAGTATCCAATTAGTGGATTGTTATCAGATGAATT C G C G C A C C A C C T A G C G	2000
2001	TTGTCTGGATGAAAAGCGAGAATTGTCGAGAAAGTCAA C C C C G T C C T	2040
2041	CATGCGAAGCGACTCAGTGAATGAGCGGAATTACTTCAAG C C T C C A C C T G	2080
2081	ATCCAAACTTCAGAGGGATCAATAGACAACCAGACCGTGG C T C A A C C G G A	2120

FIG. 14C

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2121	CTGGAGAGGAAGTACAGATATTACCATCCAAGGAGGAGAT T G T C C G G C C C	2160
2161	GACGTATTCAAAGAGAATTACGTACACTACCGGGTACCG T G G C C C T C A T T	2200
2201	TTGATGAGTGCTATCCAACGTATTTATATCAGAAAATAGA C C C T C C G C G C	2240
2241	TGAGTCGAAATTAAAAGCTTATACCCGTTATGAATTAAGA C C C C T C A G C C T	2280
2281	GGGTATATCGAAGATAGTCAGACTTAGAAATCTATTTGA C C C C T C C	2320
2321	TCCGTTACAATGCAAAACACGAAATAGTAAATGTGCCAGG A G C G G C C G C	2360
2361	CACGGGTTCTTATGGCCGTTTCAGCCAAATGCCAATC T T C C A T T C T C T	2400
2401	GGAAAGTGTGGAGAACCGAATCGATGCGCGCCACACCTTG G G T C A T	2440
2441	AATGGAATCCTGATCTAGATTGTTCTGCAGAGACGGGA G C T G C C G T C	2480
2481	AAAATGTGCACATCATTCCCATCATTCAACCTGGATATT G G C C T C T C C	2520
2521	GATGTTGGATGTACAGACTAAATGAGGACTTAGGTGTAT G T C G C C A C	2560
2561	GGGTGATATTCAAGATTAAGACGCAAGATGGCCATGCAAG C C C C C A C	2600
2601	ACTAGGGAATCTAGAGTTCTCGAAGAGAAACCATTATTA T C C T G G C	2640
2641	GGGGAAAGCACTAGCTCGTGTGAAAAGAGCGGAGAAGAAGT T T C G A	2680
2681	GGAGAGACAAACGAGAGAAACTGCAGTTGGAAACAAATAT G T C G A G T C	2720
2721	TGTTTATAAGAGGCAGGAAAGAATCTGTAGATGCTTTATTT C C G C G G C G C	2760
2761	GTAAACTCTCAATATGATAGATTACAAGTGGATACGAACA G C C A G G C C C	2800
2801	TCGCCATGATTGATGCCGGCAGATAAACGCGTTCATAGAAT C C C C T G C C C	2840

FIG. 14D

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2841	CCGGGAAGCGTATCTGCCAGAGTTGTCTGTGATTCCAGGT T T G T C T T C C T	2880
2881	GTCAATGCGGCCATTTCGAAGAATTAGAGGGACGTATTT G C T C G C T C	2920
2921	TTACAGCGTATTCTTATATGATGCGAGAAATGTCATTA C A T C G C C C C	2960
2961	AAATGGCGATTCATAATAATGGCTTATTATGCTGGAACGTG G C T C C C CAGC T	3000
3001	AAAGGTCATGTAGATGTAGAAGAGCAAAACAACCACCGTT G C G G A G T G	3040
3041	CGGTCTTGTATCCCAGAATGGGAGGCAGAAGTGTACAC C G G G T G A T C	3080
3081	AGAGGTTCGTGTCTGTCCAGGTCTGGCTATATCCTTCGT A A A A C T C	3120
3121	GTCACAGCATATAAAGAGGGATATGGAGAGGGCTGCGTA G C T C G C T T G	3160
3161	CGATCCATGAGATCGAAGACAATACAGACGAACGTAAATT C C G A C C G T G	3200
3201	CAGCAACTGTGTAGAAGAGGAAGTATATCCAAACAACACA TC C C G A A C C C	3240
3241	GTAACGTGTATAATTATACTGGGACTCAAGAAGAATATG T T C C G C T A G G C	3280
3281	AGGGTACGTACACTTCTCGTAATCAAGGATATGACGAAGC G A G C A G C T C A	3320
3321	CTATGGTAATAACCCCTTCCGTACCGACTGATTACGCTTCA TCC TCXXXXXXXXXX T T C T C C	3360
3361	GTCTATGAAGAAAAATCGTATACAGATGGACGAAGAGAGA G C G G C C C A C T	3400
3401	ATCCTTGTGAATCTAACAGAGGCTATGGGATTACACACC C C G T C T C A C	3440
3441	ACTACCGGCTGGTTATGTAACAAAGGATTTAGAGTACTTC T A T C T C G C T T	3480
3481	CCAGAGACCGATAAGGTATGGATTGAGATCGGAGAAACAG T C A G C T C	3520
3521	AAGGAACATTCATCGTGGATAGCGTGGATTACTCCTTAT G C C G C T T G	3560
3561	GGAGGAA 3567	

FIG.14E

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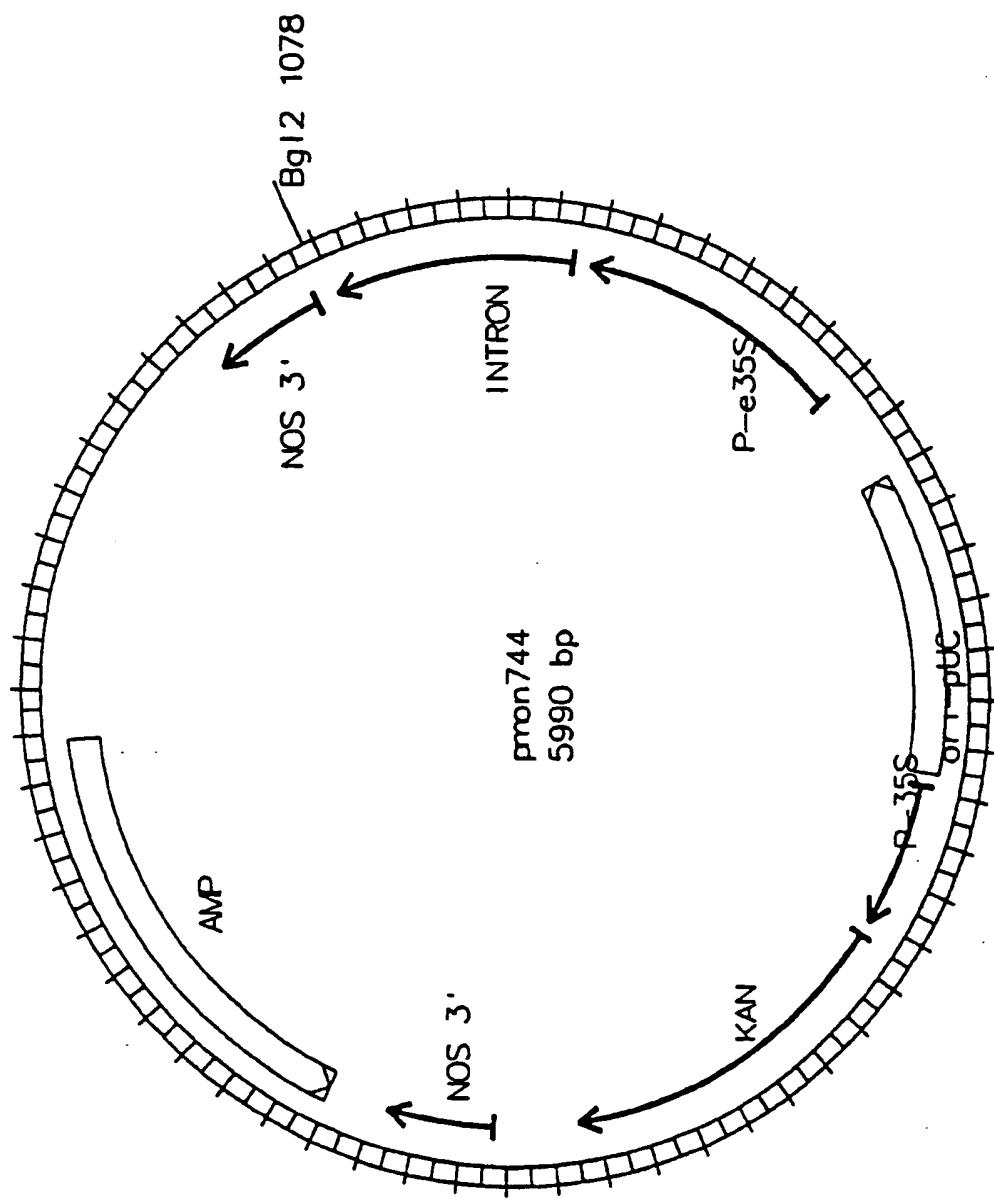


FIG. 15

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1	AGATCTAGAGGTAATTGTTATGAGTACTGTCGTGGTTAAG GATC	40
41	GGAAACGTCAACGGTGGTACAACAAACCTAGAAGGAGGA G T A	80
81	GAAGGCAATCCCTCGCAGGAGGGCTAACAGAGTACAGCC T A T	120
121	AGTGGTTATGGTCACTGCTCCTGGCGAACCCAGGAGGAGG GC A A A	160
161	AGACGCAGAAGAGGAGGCAATCGCAGGTCAAGAAGAACTG A G T A	200
201	GAGTTCCCAGGGGAAGGGGCTCAAGCGAGACATTGTGTT A A T	240
241	TACAAAGGACAACCTCGTGGCAACTCCCAAGGAAGTTTC	280
281	ACCTTCGGACCAAGTGTATCAGACTGTCCAGCATTCAAGG T	320
321	ATGGAATACTCAAGGCCTACCATGAGTACAAGATCACAAG T	360
361	TATCCTTCTTCAGTCAGCGAGGCCTTCCACCTCA T G T	400
401	CCAGGATCCATCGCTTATGAGTTGGACCCACATTGCAAAG C A T	440
441	TATCATCCCTCCAGTCCTACGTCAACAAAGTTCAAATCAC T	480
481	AAAGGGAGGAGCTAAGACCTATCAAGCTAGGATGATCAAC T T C T	520
521	GGAGTAGAATGGCACGATTCATCTGAGGATCAGTCAGGA T T A	560
561	TACTTTGAAAGGAAGTGGAAAATCTTCAGACCCAGCAGG C A G T T	600
601	ATCTTCAGAGTCACCATCAGAGTGGCTTTCAAAACCC T T A	640
641	AAGTAATAGACTCCGGATCAGAGCCTGGTCCAAGCCCACA A T	680

FIG. 16A

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681 ACCAACACCCACTCCAACCTCCCCAAAAGCATGAGCGATT 720

721 ATTGCTTACGTCGGCATACTATGCTGACCATTCAAGAAT 760

761 TC 762

FIG. 16B

# INTERNATIONAL SEARCH REPORT

International Application No. PCT/US 90/00778

## I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) \*

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC<sup>5</sup> C 12 N 15/82, 15/32, 15/67, 15/40, 5/10, A 01 H 5/00

## II. FIELDS SEARCHED

Classification System	Classification Symbols	Minimum Documentation Searched †
IPC <sup>5</sup>	C 12 N, A 01 H	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched *		

## III. DOCUMENTS CONSIDERED TO BE RELEVANT \*

Category *	Citation of Document, †† with indication, where appropriate, of the relevant passages †‡	Relevant to Claim No. †§
Y	EP, A, 0142924 (LUBRIZOL GENETICS INC.) 29 May 1985 see page 26, lines 20-30 --	1,30,31
Y	UCLA Symp. Mol. Cell. Biol., New Ser. volume 48, 1987, Molecular Strategies for Crop Protection, Alan R. Liss, Inc., M.J. Adang et al.: "Expression of a <u>Bacillus thuringiensis</u> insecticidal crystal protein gene in tobacco plants", pages 345-353 see page 351, last paragraph --	1,30,31
A	Plant Physiol., volume 85, 1987, K.A. Barton et al.: "Bacillus thuringiensis $\delta$ -endotoxin expressed in transgenic Nicotiana tabacum provides resistance to lepidopteran insects", pages 1103-1109 see page 1109, left-hand column, last two paragraphs -- --	3-28 ./.

\* Special categories of cited documents: †

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"Z" document member of the same patent family

## IV. CERTIFICATION

Date of the Actual Completion of the International Search  
5th June 1990

Date of Mailing of this International Search Report

09.07.90

International Searching Authority

EUROPEAN PATENT OFFICE

Signature of Authorized Officer

M. Peis

M. PEIS

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category	Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages	Relevant to Claim No.
O,A	Biological Abstracts/RRM, BR 35:107674, AN 88:477784, M. Adang et al.: "Engineering crop plants for insect resistance", & 154th National American Association for the Advancement of Science, Annual meeting, Boston, Massachusetts, USA, February 11-15, 1988 see the abstract	3-28
A	EP, A, 0275957 (HOECHST) 27 July 1988 see the whole document	4
A	-- Nucleic Acids Research, volume 17, no. 2, 1989, IRL Press, (Oxford, GB), E.E. Murray et al.: "Codon usage in plant genes", pages 477-498 see the whole document	4
A	-- EP, A, 0223452 (MONSANTO) 27 May 1987 see example 11	28
E	-- EP, A, 0359472 (LUBRIZOL GENETICS) 21 March 1990 see page 22, lines 1-14; claim 12	1-14, 25, 28, 30-38
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ANNEX TO THE INTERNATIONAL SEARCH REPORT  
ON INTERNATIONAL PATENT APPLICATION NO.

US 9000778  
SA 34761

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 29/06/90. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

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		AU-A-	3347384	28-03-85
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EP-A- 0359472	21-03-90	AU-A-	4118289	15-03-90

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